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This guidance applies to new content or specific content undergoing revision as part of an approved project. There are therefore many terms in the existing content (LEGACY CONTENT) that do not comply with this guidance. The process of correcting existing content will be carried out as time and resources permit and as per an approved project.

The Editorial Guide provides the information necessary to model terms in SNOMED CT. It is for those who edit content in the International Release, but it may also be useful to those creating extensions. It is a working document, subject to change and revision.

SNOMED CT is distributed in sets of electronic files. Supporting software tools are not necessarily provided directly by SNOMED International.

Short link: http://snomed.org/eg

The Editorial Guide publishes only the rules that apply to precoordinated content. That is, the rules where the 'Content Type' is one of the following - [All SNOMED CT content], [All precoordinated SNOMED CT content], or [All new precoordinated SNOMED CT content]. The rules that are not published in the Editorial Guide are the ones that apply only to [All postcoordinated content]. See the different content types and rules in the MRCM maintenance tool at https://browser.ihtsdotools.org/mrcm.

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SNOMED CT Introduction

What is SNOMED CT?

SNOMED CT is a high-quality, comprehensive, international, logic-based reference terminology that is used to present clinically relevant information. It began with the union of NHS Clinical Terms Version 3 and SNOMED RT;
this provided the initial scope which has since been updated to reflect contemporary clinical practice and changes in medical technology.

Content development is provided by expert clinicians driven by the requirements of user communities. This includes core content for use internationally and content relevant to national extensions for local implementation.

Its logic-based definitions represent terminological knowledge, or what is always true about the meaning of concepts. It consists of codes, that correspond to concepts, arranged in a polyhierarchical manner, as well as relationships between the concepts, which further define the meaning.

Why use SNOMED CT?

It supports semantic interoperability and multi-purpose use within electronic health applications (primarily electronic health records or EHRs) and has many advantages over other terminologies. They include:

- Consistent, and formal expansion of, content through centralized authoring and maintenance (International Release)
- Flexibility to meet most terminological requirements based on national, regional, language, application, or customer (Extensions)
- Clear, singular meaning of concepts
- Reliable, consistent, and reproducible clinical documentation
- Enhanced high-quality healthcare delivery to individuals and populations

Intended Use

SNOMED CT is intended to be used in healthcare:

- To provide effective and comprehensive coverage of terms
- As a terminological resource
- For implementation in electronic health applications

The purpose of SNOMED CT is to represent clinically relevant information reliably and reproducibly in electronic health applications, (most often electronic health records or EHRs) to support:

- Delivery of multidisciplinary, high-quality healthcare to individuals and populations
- Optimal retrieval, processing, and rendering of clinical information
- Effective use of clinical information consistently and reproducibly
- Use of clinical information for statistical and reporting purposes

Semantic Interoperability

The overall semantic interoperability of electronic health applications is achieved through the combined functioning of the information architecture of the application and the terminology that populates it. A basic principle of SNOMED CT is to create and maintain semantic interoperability of clinical information. Semantic interoperability is the capability of two or more systems to communicate and exchange information. Each system should be able to interpret the meaning of, and effectively use, received information. To achieve this goal, the meaning of the information must be agreed upon, consistent, and clearly expressed.
Structure of Domain Coverage

SNOMED CT includes 19 domains arranged in a polyhierarchical structure. Each hierarchy is an ordered organization of concepts linked together through IS-A relationships. Each concept may have one or more parents.

The hierarchical arrangement is helpful for locating concepts, grouping similar concepts, and conveying meaning. For example, if we see the concept cell under the concept anatomic entity we will understand the intended meaning as different than if it appeared under the concepts room or power source (Desiderata for Controlled Medical Vocabularies in the Twenty-First Century by J.J. Cimino published in Methods of Information in Medicine 1998:37:394-403).

Concepts are linked to their more general parent concept codes directly above them in a hierarchy. Concepts with more general meanings are usually presented as being at the top of the hierarchy and then at each level down the hierarchy, the meanings become increasingly more specific or specialized.

The domains contain all of the components (clinical, administrative, database structure, as well as other components that express how the domains relate to each other) necessary to create SNOMED CT concepts and maintain the database structure.
A domain is a set of concepts which the Concept Model permits to be defined or refined, using a particular set of attributes and ranges. Some domains do not have attributes and ranges but may if a concept model is created.

A domain, to which an attribute can be applied, is typically defined to include concepts in one or more branches of the subtype hierarchy.

The domain of 116676008 | Associated morphology (attribute) is defined as subtype of 404684003 | Clinical finding (finding).

The range of values of 116676008 | Associated morphology (attribute) is subtypes of 49755003 | Morphologically abnormal structure (morphologic abnormality).

The following table lists the domains, definitions, and examples. *Those without a concept model are marked with an asterisk.

<table>
<thead>
<tr>
<th>Domains</th>
<th>Examples</th>
</tr>
</thead>
</table>
| **1 Body Structure** | • Anatomical or acquired body structure  
• Morphologic abnormality (subtype of body structure) |
| | • 450807008 | Entire back (body structure) |
| | • 52988006 | Lesion (morphologic abnormality) |
| **2 Clinical Finding** | • Clinical finding: normal/abnormal observations, judgments, or assessments of patients  
• Disorder: always and necessarily an abnormal clinical state |
| | • 39579001 | Anaphylaxis (disorder) |
| | • 167222005 | Abnormal urinalysis (finding) |
| **3 Environment and Geographical Location** | • Environment: types of environments  
• Geographical Location: named locations such as countries, states, or regions |
<p>| | • 405607001 | Ambulatory surgery center (environment) |
| | • 223581004 | China (geographic location) |
| <strong>4 Event</strong> | • Occurrences impacting health or health care; not procedures or interventions |
| | • 242039002 | Abuse of partner (event) |
| | • 2641000119104 | Exposure to chlamydia (event) |
| <strong>5 Observable Entity</strong> | • Information about a quality/property to be observed and how it will be observed |
| | • 423493009 | Age at diagnosis (observable entity) |
| | • 416125006 | Concentration of hemoglobin in erythrocyte (observable entity) |
| <strong>6 Organism</strong> | • Organisms of significance to human and animal medicine; use in modeling cause of disease |
| | • 3265006 | Genus Candida (organism) |
| | • 710877000 | Beta lactam resistant bacteria (organism) |</p>
<table>
<thead>
<tr>
<th>Domains</th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>7</td>
<td>Pharmaceutical/Biological Product</td>
<td>• Drug products (not Substances)</td>
</tr>
<tr>
<td>8</td>
<td>Physical Force*</td>
<td>• Forces applied to the body that may cause injury</td>
</tr>
<tr>
<td>9</td>
<td>Physical Object*</td>
<td>• Physical devices relevant to health care, or to injuries/accidents</td>
</tr>
<tr>
<td>10</td>
<td>Procedure</td>
<td>• Procedure: activities performed in the provision of health care (includes medical history-taking, physical examination, diagnostic and therapeutic interventions, training and education, and counseling)&lt;br&gt;• Regime/therapy (subtype of procedure): set of procedures focused on a single purpose on one patient over time (e.g. repeated administration of drug in a small dose for an indefinite period of time)</td>
</tr>
<tr>
<td>11</td>
<td>Qualifier Value*</td>
<td>• One of several possible values for an attribute used to define concepts</td>
</tr>
<tr>
<td>12</td>
<td>Record Artifact*</td>
<td>• Clinical documents, or parts thereof</td>
</tr>
<tr>
<td>13</td>
<td>Situation with Explicit Context</td>
<td>• Concepts that include context information; a subtype of the situation to which it applies with an attribute associating it with the relevant clinical finding or procedure&lt;br&gt;• May be used to represent conditions/procedures that already occurred, haven’t yet occurred, or refer to someone else (not patients)</td>
</tr>
<tr>
<td>Domains</td>
<td></td>
<td></td>
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<tr>
<td>---------</td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>14. SNOMED CT Model Component</strong>&lt;sup&gt;*&lt;/sup&gt;</td>
<td>• Concepts and attributes necessary to organize and structure SNOMED CT terminology and its derivatives</td>
<td>• 900000000000442005</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• 900000000000454005</td>
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<tr>
<td></td>
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<td>• 106237007</td>
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<td></td>
<td>• 370136006</td>
</tr>
<tr>
<td><strong>15. Social Context</strong>&lt;sup&gt;*&lt;/sup&gt;</td>
<td>• Social conditions and circumstances related to healthcare</td>
<td>• 116060000</td>
</tr>
<tr>
<td></td>
<td>• Subtypes include: ethnic group, life style, occupation, person, racial group, religion/philosophy, social concept</td>
<td>• 58626002</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• 415794004</td>
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<tr>
<td></td>
<td></td>
<td>• 35359004</td>
</tr>
<tr>
<td><strong>16. Special Concept</strong>&lt;sup&gt;*&lt;/sup&gt;</td>
<td>• Inactive and navigational (support locating concepts in hierarchies) concept codes</td>
<td>• 363664003</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• 394899003</td>
</tr>
<tr>
<td><strong>17. Specimen</strong></td>
<td>• Entities that are obtained (usually from patients) for examination or analysis</td>
<td>• 373193000</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• 258441009</td>
</tr>
<tr>
<td><strong>18. Staging and Scales</strong>&lt;sup&gt;*&lt;/sup&gt;</td>
<td>• Assessment and tumor staging scales</td>
<td>• 273472005</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• 254294008</td>
</tr>
<tr>
<td><strong>19. Substance</strong></td>
<td>• Active chemical constituents of allergens, agents, substances, chemicals, drugs, and materials (not Pharmaceutical/Biological Products)</td>
<td>• 116272000</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• 64856004</td>
</tr>
</tbody>
</table>

**Granularity**

The scale, or level of detail, in a terminology is called *granularity*. Concepts and meanings range from very general, or coarse, to very specific, or fine. SNOMED CT has multiple granularities, which is an important component of
terminologies that are multipurpose. The broader meanings are useful for aggregation (e.g. Clinical finding, Procedure, etc.), but are not intended for recording individual patient data.

The progressive levels of refinement are used to meet clinical data requirements. There are, however, limits to the degree of precoordination of certain types of complex statements.

In general, concepts in SNOMED CT should name things that exist in the real world. The concepts are usually names or short noun phrases, not complete sentences or paragraphs.

SNOMED CT is intended to be used with electronic health applications that can support full clinical statements, along with their attributes, dates, times, and statement interrelationships. It may be challenging to balance SNOMED CT content with the needs of those using electronic health applications. For example, some older applications may require concepts outside of the scope of SNOMED CT. SNOMED CT tries to maximize its usefulness and at the same time minimize precoordination.

Knowledge Representation

Knowledge representation in SNOMED CT involves modeling what we know about concepts to be necessarily true. Concepts are logically defined by their relationships to each other. Some knowledge provides valuable clues to the diagnostician, while not necessarily always present, i.e. it is uncertain or probabilistic knowledge. Attempts to capture probabilistic or uncertain knowledge are out of the scope of SNOMED CT.

For example,

- 22298006 | Myocardial infarction (disorder) |
  Its terminological knowledge includes the following:
  - IS A: 64572001 | Disease (disorder) |
  - Finding site: 74281007 | Myocardium structure (body structure) |
  - Associated morphology: 55641003 | Infarct (morphologic abnormality) |

  These additional pieces of knowledge are variably present and therefore represent uncertain or probabilistic knowledge about myocardial infarction:
  - Crushing substernal chest pain
  - Diaphoresis
  - Arrhythmia
  - ST-segment elevation on EKG
  - Elevated cardiac enzymes

For example,

- 74400008 | Appendicitis (disorder) |
  Its terminological knowledge includes the following:
  - IS A: 64572001 | Disease (disorder) |
  - Finding site: 66754008 | Appendix structure (body structure) |
  - Associated morphology: 23583003 | Inflammation (morphologic abnormality) |

  These additional pieces of knowledge are variably present and therefore represent uncertain or probabilistic knowledge about appendicitis:
  - Central abdominal pain that migrates to the right lower quadrant
  - Rebound tenderness over McBurneys point
  - Anorexia
  - Nausea
  - Elevated white blood count
Out of Scope

National and local extensions

SNOMED CT has an international and multilingual scope but can be localized to represent meanings and terms unique to particular organizations or localities. A National Extension includes content outside of the scope of the International Release, but necessary for national conformance and interoperability. Each member-state determines the application and interpretation of this scope and whether or not concepts should be added to their extension.

National Extension criteria include affirmative answers to the following:

- Is the concept outside of the scope of the International Release, but necessary for national conformance and interoperability?
- Is it useful throughout the national healthcare system?
- Does it need to be understandable throughout the national healthcare system?
- Does it need to be shared in a reproducible manner within the national healthcare system?

Extensions are created, structured, maintained, and distributed in accordance with SNOMED CT specifications and guidelines to ensure compatibility with the SNOMED CT International Release. Members may create, maintain, and distribute extensions to address specific national, regional, and language requirements. Affiliates may also create, maintain, and distribute extensions to meet the needs of particular software solutions and customers. Content that is within the scope of the International Release is restricted to the International Release and may not be modified or replaced by an extension, unless explicitly permitted by SNOMED International. Please see the Practical Guide to Extensions for more information.

Veterinary extension

SNOMED CT is not intended to cover all medical knowledge. Content that is strictly related to animals is out of the scope of the SNOMED CT international release. Non-human content may be included in a request for new content via the SNOMED International Content Request System (CRS) or may be identified in the International Release. Careful consideration is required to differentiate content that belongs in the International Release versus an extension. The basic principle is that content used in human medicine should be in the core. Content that is strictly non-human may be managed in an extension.

Examples of non-human content,

- Egg-related coelomitis (disorder)
- Dehorning (procedure)
- Bone structure of wing (body structure)

Types of content that should be in the core include the following:

- **Diseases and findings.** Anything that can occur in both humans and animals should be in the core.
• **Material entities.** A material entity is a concept found within the Substance, Physical object, Pharmaceutical/biologic product, Physical force, or Organism subhierarchies. Every substance that can cause adverse effects should be in the core (with the understanding that poisonings and adverse effects in humans may be caused by virtually any substance). Some material entities may be of interest only in a non-human or veterinary context. These entities may be added to, or left in, a veterinary extension.

• **Organisms.** Most organisms should be in the core, with some exceptions. There are almost 27,000 organism codes in the Veterinary Extension maintained by the Veterinary Terminology Services Laboratory (VTSL) at Virginia Tech University. Generally, most of these are descendants of Kingdom Animalia. These are not transferred to the core, except when used in public health or human medicine or when requested by more than one SNOMED International member country. Breeds are primarily restricted to the veterinary domain. In addition, requests for other organisms that are not used in public health or human medicine can be added to the Veterinary Extension. This is mostly applicable to macroorganisms as microorganisms can change host or take advantage of immunosuppression in humans. Also, human laboratories may need to report animal pathogens. Organisms that are not used in human medicine can be added to the Veterinary Extension.

The Veterinary Extension is publicly available to SNOMED International member countries and to Affiliate Licensees. To access to the Veterinary Extension, please see [http://vtsl.vetmed.vet.edu](http://vtsl.vetmed.vet.edu), or contact the VTSL at vtsl.extension@gmail.com.

**Classification-derived phrases**

Classification-derived phrases are not accepted. Concepts with unclear, unspecified, or ambiguous meaning should not be used. Rejections are expected for requests with the following phrases:

- Not otherwise specified (NOS)
  - For example, Mental disorder, *not otherwise specified*
- Not elsewhere classified (NEC)
  - For example, Chronic hepatitis, *not elsewhere classified*
- No mention
  - For example, Bile duct calculus with *no mention* of cholecystitis and with obstruction
- With or without
  - For example, Tubal pregnancy *with or without* intrauterine pregnancy

**Regulatory status or characterization**

Concepts referring to regulatory status or characterization (e.g., over-the-counter) are out of scope for the International Release. Meaning may vary by jurisdiction and may not be consistent internationally.

**SNOMED CT Requirements**

Key requirements that drive the design, development, and maintenance of SNOMED CT are as follows. They are related to:

1. Electronic health applications (most often electronic health records or EHRs)
   - Support for effective delivery of high quality healthcare to individuals and populations
2. The terminology
3. Implementation and migration
4. The intended user communities
   - International, multilingual applicability
   - Supporting particular localities
5. National and strategic priorities

These requirements are interrelated. The design objective is to enable all user communities to realize the potential benefits. However, the needs of different user communities may vary. To meet the overall objectives, the design must consider the entire range of needs. The approach must also be scalable in order to enable extension to new user communities.
Medical Vocabularies - J. Cimino

The headings in this section are the requirements identified in Desiderata for Controlled Medical Vocabularies in the Twenty-First Century by J.J. Cimino published in Methods of Information in Medicine 1998:37:394-403. Following each, is an explanation of the way in which SNOMED CT meets the requirement.

Content, content, and content

SNOMED CT content must be adequate both in scope and quality and must:

- Cover a wide variety of domains and different organizational needs, clinical disciplines, and medical specialties
- Meet the needs of an expanding scope, while retaining quality, with a structured systematic approach

Nonvagueness and nonambiguity

Codes must have one meaning (nonvagueness) and no more than one meaning (nonambiguity). These characteristics are sometimes called concept orientation, but SNOMED CT deprecates the use of the word concept to describe codes or their meanings.

A code and its meaning may be expressed by more than one term. The terms vary between languages and dialects. In any language or dialect there may be several synonymous terms.

Code permanence

Once assigned a meaning, a code must not change its meaning. Refinements, due to changes in the state of knowledge, may lead to inactivation of codes from SNOMED CT. An inactivated code may be replaced by a new, more precisely defined code.

Nonsemantic identifiers

The structure of an identifier (code) should not contain any semantic information about its meaning or relationships.

Polyhierarchy

SNOMED CT supports multiple hierarchies. A code may have more than one hierarchical parent and various paths to its root code.

Formal definitions

When possible, the meaning of codes should be formally defined by relationships to other codes.

Rejection of Not elsewhere classified

Codes with the phrase, not elsewhere classified, are not allowed in SNOMED CT. However, many classifications contain terms with this phrase. A term with not elsewhere classified includes general variants that are not specifically represented. The meaning of such a code may change over time. As codes with more specific meanings are added, this narrows the codes included in the not elsewhere classified codes.

Multiple granularities

Different users will need to express more or less finely granular meanings. SNOMED CT:

- Must accommodate a wide range of levels of detail
- Must recognize the relationships between meanings at different levels of granularity
- Should allow selection of codes that include navigation to other codes with more or less finely grained meaning
• May need to restrict the levels of granularity used in different applications or in different contexts within the same application

Multiple consistent views
The view of a code's meaning, with multiple hierarchical parents, should not depend on reaching it by following the hierarchy from a particular parent.

Beyond terminology codes - represent context
The meaning of a code in a patient record may be altered by its context. Standards for patient record architectures and modeled healthcare communication are changing. The role of SNOMED CT in the context of these structures should be evaluated and appropriate recommendations made.

Evolve gracefully
Terminologies need to change over time. SNOMED CT should implement these changes in ways that are well-documented and tracked and that provide a path for systems and users.

Recognize redundancy
The same information can often be coded in different ways. A controlled terminology, that has an adequate scope, cannot exclude this possibility. Instead it should facilitate recognition of equivalent terms.

Electronic Health Applications
The anticipated benefits of SNOMED CT are derived from use of information to support effective delivery of high quality healthcare to individuals and populations.

Individuals

Aide-memoire for clinicians
Clinically relevant information in an electronic health record acts as an aide-memoire for the clinician, enabling recall of previous interactions.

Structured data entry
Structured data entry enhances the value of an electronic health record in various ways. It may:

• Simplify recording of frequently collected data
• Ensure that information is collected in a reliable and reproducible way
• Help clinicians to think logically about a patient's condition

Clinical applications may combine several data entry methods. Some of the most commonly used methods are as follows:

• Searching a coded terminology for matching terms using words or phrases
• Navigating a hierarchical structure to refine or generalize meanings
• Using templates or protocols to record structured information; may be based on answers to questions or values entered on a data entry form
• Parsing of natural language to identify and retrospectively code and structure data
• Typing, speech recognition, and document scanning

SNOMED CT requirements for data entry
Data entry may require selection from a list. Such lists must be manageable in size and appropriate to the needs of the user.
A multilingual, multidisciplinary terminology requires mechanisms that limit and/or prioritize access to
terms and codes in ways that are appropriate to:
  • Languages and dialects
  • Countries, organizations, disciplines, specialties, and users
  • Contexts within a record or protocol

To display a code’s description in a list that has not been derived from a text search, the term must be
intelligible and appropriate to the user.

When a code is entered in a record it may require structured entry of additional qualifying information.
  • Qualifying information may be coded.
    For example, the code named *removal of kidney* may require a statement of laterality.
  • Qualifying information may be numeric.
    For example, the code named *hemoglobin measurement* may enable entry of a numeric value expressed
    in a substance concentration.

To meet all the needs for coded structured data entry in a health record, a terminology must have an adequate
scope.
  • The main body of SNOMED CT covers the required scope.
    • It may be difficult to meet the needs of some organizations, specialties, and users; they may need
      specific terms or codes to meet their own operational requirements. Therefore, SNOMED CT is
      structured to allow for additions to meet specific needs.

A clinical terminology requires frequent changes including new codes, terms, and relationships between codes.
Changes may be required due to new:
  • Health risks
  • Health and disease process information
  • Drugs, investigations, therapies, and procedures

**Presentation**

The presentation of clinical information may:
  • Highlight key information and indicate links between items, thus helping clinicians understand patients' conditions.
  • Be determined entirely by record structure without regard to the terminological resource (e.g., may be in
    chronological order, by author, or by the type of recorded event).
  • Be enhanced based on its semantic content (e.g., grouping procedures, investigation results, or observations relevant to a particular disease process).

**Decision support**

Interfaces between recorded clinical information and appropriate decision support tools and reference works may assist the clinician in selecting diagnostic tests, making diagnoses, and choosing treatment. Decision support requires selective retrieval and processing of information in an individual health record to determine whether the patient has particular characteristics relevant to the decision support protocol. The algorithms for establishing the presence of characteristics should include relationships between coded meanings and other aspects of record structure. Performance is also important, as decision support algorithms are typically run in real-time during data recording. Decision support algorithms may:
  • Depend on numeric or other values (and their units) associated with particular observations
  • Include the context in which information is recorded, e.g., the date of recording and any stated
    relationships between individual items of information
  • Include information such as age, sex, clinical conditions, findings, surgical procedures, medication, and
    social/environmental factors, such as occupation
  • Use codes or identifiers from other terminologies, classifications, or proprietary schemes. Mapping tables are
    required to allow applications that use a terminology to interface with these resources
Communication

Effective delivery of high quality healthcare to individuals requires communication between those involved in providing care. This requires communication within and across teams or organizations.

The primary objective of many clinical communications is to convey information from human to human. Communications with this purpose should include human-readable text. Relying on text from coded data is not recommended. Coded data is therefore not relevant to the requirement for human-to-human communication.

A receiving application may process clinical communications. This information may need to be retrieved and processed to meet terminology requirements. To meet terminology requirements, messages and other means of electronic communication must permit the communication of SNOMED CT identifiers and associated structures.

Communication specifications, such as those produced by HL7 and CENTC251, define the structures to meet requirements. The coded information is used in two distinct situations:

- Coded elements that must be filled with codes enumerated in the specifications. The codes enumerated in the specifications generally communicate, mission critical features of the message. Some of the enumerated codes and the codes in a clinical terminology may have overlapping meanings.
- Coded elements that are populated with clinical codes from appropriate coding schemes. The open coded elements may require the full expressiveness of a terminology. Some of the open coded elements may be restricted to codes that express particular types of meaning.

For example, HL7 requires that coding schemes meet certain criteria, one of which is the ability to express limited subsets of codes appropriate to particular elements.

There are two situations in which communication of coded information may be of value for human-to-human communication. They are where:

- The storage capacity or communication bandwidth is restricted. Receiving applications must contain (or have real-time access to) a table listing the text description associated with each code.
- The translation between the languages of the sender and the recipient is needed. A coded representation of a meaning may allow the appropriate description in the recipient’s language.

Recording a particular code may trigger a communication. And, receipt of a code, may trigger specific processing in the receiving application.

For example, recording a decision to prescribe a medicine might trigger an electronic prescription sent to the pharmacy. Receipt of such a prescription might trigger dispensing and stocking activities.

The relationship of a trigger, is an additional characteristic of a code, that may be context dependent.

Patient involvement

Patients may wish to view, and comprehend, their own records. For SNOMED CT to meet this requirement, the inclusion of patient-friendly terms should be considered. However, this requirement should not take precedence over accurate professional terminology.

Patients may also be allowed to contribute to their own records, i.e. be users of SNOMED CT.

For example, patients with diabetes may monitor and record their blood glucose levels.

Populations

Identify and monitor health needs

The provision of effective high-quality care to populations requires an understanding of the state of health and healthcare needs of that population. Information recorded about individual patients must be available for analysis to determine trends.

- It must be possible to analyze data recorded with SNOMED CT.
Population trends are usually monitored at a higher level, using codes that are more general than those used in individual patient records. This may be accomplished through one or both of the following methods:

- Using hierarchical relationships and/or equivalences defined within SNOMED CT.
- Mapping SNOMED CT codes to codes in appropriate classifications.

Appropriate analysis of information requires reliable and reproducible queries.

- The scope of SNOMED CT must cover the types of information relevant to analysis.
- Analysis may require data about multiple clinical characteristics. Queries must account for both the terminology and the record structure.

Audit quality of service

The requirements for analysis of quality of service are similar to those for analysis of health needs. The main difference is that the scope of the analysis must be extended to cover consultations, referrals, procedures, medications, and other interventions.

Support research

The requirements for research are also similar to those for analysis of health needs, however, there is a need to allow for:

- Recording interventions in ways that do not compromise blind and double blind trials.
- Adding SNOMED CT content for experimental observations or treatments, which may never require permanent addition to the terminology.

Reduce bureaucracy; manage and fund care delivery

The management and funding of healthcare delivery often depends on recording and reporting of particular information, e.g. bundled or packaged care. Automating this process offers a way of reducing bureaucratic overhead, i.e. mapping clinical information recorded with SNOMED CT to appropriate forms.

Some information required for management and funding purposes is specifically related to claims for particular events or services. For example, funding general practitioners in the NHS is dependent on meeting immunization administration and cervical cytology screening targets.

The scope of SNOMED CT must be adequate to meet these needs, or must be capable of extension to meet these needs, without presenting irrelevant terms or coded meanings to those not requiring them.

Enable reporting of external health statistics

Organizations, such as WHO and some government bodies, require specific data related to healthcare statistics. Organizations should be able to use clinical information recorded with SNOMED CT. When this is not possible, the clinical information should at least support their manual generation. Using structured data entry allows for direct mapping to statutory national and international classifications such as ICD, CPT, OPC, and etc.

Identify patients in need of interventions proactively

Population-based preventive care should be offered to specific groups, based on sex, age, medical history and other factors. Health information applications based on information recorded with SNOMED CT can be used to identify patients so they can be offered appropriate care.
Implementation and Migration

Electronic health application
A terminological resource is only one part of an electronic health application. Implementation of SNOMED CT should support applications in meeting user needs, rather than adding a burden to development.

The functions required to implement a terminology can be divided into those that are:

- Performed without reference to data stored in a particular application record structure.
- Involved in storing, retrieving, or processing application data.

Applications may make use of different aspects of SNOMED CT. Some may require SNOMED CT for a very limited range of uses for which there may be minimal value. These applications may not require all the functions for a full implementation or all the concepts and codes in SNOMED CT.

- There may be a general benefit in consistency with other more terminology rich applications.

Existing information
A substantial body of clinical information may already be present in an electronic health application. Much of this information is represented using existing coding schemes, terminologies, and classifications. This information may be of value to individual patient records or to populations. Similarly, there are many queries and decision support protocols that contain information based on existing terminologies.

A new terminology should make provisions for the continuing use of information stored in records, queries, and protocols represented by other terminologies. There are two general approaches to this:

- Conversion of legacy data into a form consistent with SNOMED CT.
- Allowing legacy and SNOMED CT data to coexist. Legacy codes must be recognizably different from SNOMED CT codes. In addition, the relationship between codes in SNOMED CT and legacy codes must be recognized when retrieving data.

Reliability and reproducibility
Information represented with SNOMED CT codes must be reliable and reproducible. This means:

- The meaning of a code should not change over time.
- Information should be reproducible independent of the application.
- The query of codes should be reliable. This means:
  - There should be complete recall, including specific, more detailed codes and expressions subsumed by general codes and expressions in the query.
  - There should be specificity and precision excluding codes and expressions that are not subsumed by the codes and expressions in the query.
  - The effects of the following should be taken into account:
    - Precoordinated relationships between codes in records or queries.
    - Postcoordinated qualifications applied to codes or expressions in records or queries.
    - Relationships between codes and other contextual information implied by the record structure.

User Communities

Language
The terms required by users of a clinical terminology vary according to the local languages and dialects.

- When using a terminology, users must see terms in a language and dialect with which they are familiar. The terms must be clear and unambiguous independent of any hierarchical context or formal definition.
- The display of terms must not be confused by inclusion of terms in other languages or dialects.
The terms used in different languages and dialects are not mutually exclusive. A term may be common to several languages or dialects of a language.

When a code is presented without a specific reference to a term, an appropriate preferred term should be displayed. A term may be a preferred term in one dialect and a synonym in another.

Some terms differ only in spelling conventions (e.g. color vs. colour). The same spelling variants may recur in many different terms.

It may be appropriate to recognize these cases and handle them differently from other term variants.

An individual instantiation of an application may only require access to a single language or dialect. It is inappropriate to install and maintain all language and dialect variants.

An application may need to support several languages with the ability to switch between languages and dialects in real-time to meet the needs of users.

**Specialty**

Some specialties or disciplines prefer to use different terms to describe the same meaning. A particular specialist may use a precise term, while a generalist may use a different term to describe the same condition.

**Use of terms**

The following table lists factors affecting term use and examples of each.

<table>
<thead>
<tr>
<th>Factors affecting term use</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Factor</td>
<td></td>
</tr>
<tr>
<td>Geographic and seasonal differences</td>
<td>Malaria is more common in certain regions</td>
</tr>
<tr>
<td></td>
<td>Hay fever is more common in spring, summer, and fall</td>
</tr>
<tr>
<td>Cultural perceptions of health</td>
<td>Acceptance of alternative therapies</td>
</tr>
<tr>
<td>Discipline or specialty</td>
<td>Obstetricians use fundus to mean fundus of the uterus; gastroenterologists use the same term to mean fundus of the stomach</td>
</tr>
<tr>
<td></td>
<td>Surgeons record operative procedures relevant to their specialties</td>
</tr>
<tr>
<td>Professional criteria</td>
<td>The definition of hypertension may vary based on professional guidelines</td>
</tr>
<tr>
<td>National or organizational requirements, including those for administrative or funding purposes</td>
<td>Performance measure results affecting reimbursement</td>
</tr>
<tr>
<td>Topics of special interest to individual clinicians</td>
<td>Infectious disease specialist with an interest in tropical diseases</td>
</tr>
</tbody>
</table>

**Organization, country, and user**

Particular terms may be specific to an organization. They may not be included in the International Release of SNOMED CT. Organizations and users must be able to add terms or codes to SNOMED CT, without devaluing the main body of SNOMED CT.

It may be necessary to combine several subsets and extensions to meet the needs of a country, an organization, or a specialty. There must be consistent rules for combining subsets and extensions.

The requirements of a particular user may change according to the role they are performing. A single instance of an application may need to support different requirements of several users.
Summary of SNOMED CT Requirements

A summary of the SNOMED CT requirements is as follows. Additional information may be found throughout this guide, as well as in other documents on the SNOMED International website at: http://www.snomed.org/snomed-ct/learn-more.

<table>
<thead>
<tr>
<th>Terminology Structure</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Coded meaning</strong></td>
<td>• The central component is coded meanings</td>
</tr>
<tr>
<td></td>
<td>• Each code must have a single clear and unambiguous meaning</td>
</tr>
<tr>
<td><strong>Identifier</strong></td>
<td>• Components must have unique identifiers</td>
</tr>
<tr>
<td></td>
<td>• The internal structure of these identifiers must not imply the meaning or relationships of a code</td>
</tr>
<tr>
<td><strong>Description</strong></td>
<td>• Represents the association between terms (text strings) and the meanings that they describe (may be language or dialect dependent)</td>
</tr>
<tr>
<td><strong>Preferred Term</strong></td>
<td>• Represents the special association between each code and a preferred term (used to display the meaning, unless there is an alternative preference)</td>
</tr>
<tr>
<td></td>
<td>• The preferred term association is language or dialect dependent</td>
</tr>
<tr>
<td><strong>Fully Specified Name</strong></td>
<td>• Provides each code with a structured fully specified name that unambiguously describes its meaning</td>
</tr>
<tr>
<td></td>
<td>• The fully specified name is defined in a reference language (the language of first use)</td>
</tr>
<tr>
<td></td>
<td>• Translations of the fully specified name may also be required</td>
</tr>
<tr>
<td><strong>Hierarchy</strong></td>
<td>• Represents hierarchical relationships between coded meanings</td>
</tr>
<tr>
<td></td>
<td>• The form of representation allows a coded meaning to have multiple hierarchical parents (supertypes)</td>
</tr>
<tr>
<td></td>
<td>• It guarantees that any alternative hierarchical view of a coded meaning is consistent</td>
</tr>
<tr>
<td><strong>Relationship</strong></td>
<td>• Represents non-hierarchical relationships between coded meanings</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Content</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Scope</strong></td>
<td>• The scope is adequate to meet the requirements of various countries, organizations, disciplines, and specialties</td>
</tr>
<tr>
<td></td>
<td>• The extent to which the content requirements are covered develops over time</td>
</tr>
<tr>
<td></td>
<td>• However, the initial release should cover:</td>
</tr>
<tr>
<td></td>
<td>• The scope of the existing clinical terminologies</td>
</tr>
<tr>
<td></td>
<td>• All versions of the Read Codes and NHS Clinical Terms</td>
</tr>
<tr>
<td></td>
<td>• All versions</td>
</tr>
<tr>
<td></td>
<td>• Other scope requirements identified by the Editorial Board</td>
</tr>
<tr>
<td><strong>Updates</strong></td>
<td>• The content is regularly updated</td>
</tr>
<tr>
<td><strong>Granularity</strong></td>
<td>• Allows coded meanings to be expressed at different levels of granularity</td>
</tr>
<tr>
<td><strong>Not Elsewhere Classified (NEC); Not Otherwise Specified (NOS)</strong></td>
<td>• Codes with not elsewhere classified or not otherwise specified must be inactivated and no new ones may be added</td>
</tr>
</tbody>
</table>
### Content

| Extension         | • Allows extensions to the main body of work  
|                  | • Extensions are distinguishable from components of the main body; should be traceable to a responsible organization  
|                  | • Allows for distinguishing and tracing the code source or identifier used in patient records |

### Maintenance and Distribution

| Distribution       | • Distributed in a format that is readily usable by application developers  
|                   | • This format is fully specified and is not changed from release to release  
|                   | • May be distributed for use with associated software, such as a browser |
| Persistence        | • The meaning of a code is persistent; It is not changed or deleted by updates  
|                   | • A code may be marked as inactivated when its meaning is found to be ambiguous, redundant or otherwise incorrect  
|                   | • Changes to the association between a concept and a code do not change or delete the description. The description is marked as inactivated, and a new corrected description is created |
| History            | • All changes to components are tracked and saved in history files (includes details about new components and changes to the status of components)  
|                   | • When a component is made inactive, relationships or references indicate the replacement or equivalent component |

### Subsets

| Concepts | • Includes a mechanism for representing subsets of concepts appropriate for a language, dialect, or specialty. It should allow:  
|          | ▪ Specification of the synonyms, preferred terms, and translated fully specified names in each language or dialect  
|          | ▪ Rational combination of languages and modification of language subsets to meet the needs of organizations or specialties |
| Codes    | • Includes mechanisms for representing subsets of codes for a country, organization, discipline, or specialty. The form of representation should allow:  
|          | ▪ An indication of the priority, or frequency of use  
|          | ▪ Rational combinations of subsets to meet the needs of users or groups of users |
| Specified Contexts | • Includes mechanisms for representing subsets of codes and concepts for particular contexts in a record, decision support protocol, or data entry field |
| Combinations | • Include consistent rules for combining subsets to meet the requirements of users |
### Subsets

| Distribution and Installation | • Subsets are distributed in a format that is readily usable by system developers. The format is fully specified and does not vary from release to release. The distribution format allows:  
  • Subsets to be installed separately  
  • Related or interdependent subsets to be selected and installed as groups  
  • Subsets to be updated with each new release |
| Configuration | • It is possible to configure an application to use a particular subset or combination of subsets; changing configurations does not require reinstallation |

### Relationships

| Navigating Relationships | • Includes relationships that allow hierarchical navigation from a chosen code to a code that represents either a subtype or part of the chosen code  
  • Supports navigation from a specific code to more general codes that represent a supertype of that code  
  • Navigational concepts are not supported by SNOMED International |
| Aggregation of Related Codes | • Includes relationships that allow aggregation of related codes to enable comprehensive and accurate retrieval from patient records  
  • These relationships, together with appropriate history and cross-reference tables, enable the aggregation to include inactivated codes with similar or equivalent meanings |
| Defining Characteristics | • Includes formal definitions of codes represented by relationships with defining characteristics (e.g. the anatomical site of the code named *appendicitis* is the *vermiform appendix*) |
| Qualifying Characteristics | • Enables a code recorded in a patient record to be qualified by adding relevant qualifying characteristics  
  • Each qualifying characteristic is itself a code with a specified relationship to a qualified code  
  • Specifies possible qualifying characteristics for each code or for a group of related codes (e.g. an anatomical site could be added to the code named *osteoarthrosis*) |
| Kind-of-Value | • Enables codes to be qualified by the addition of relevant values  
  • Specifies the types of values that can be added to particular codes (e.g. a substance concentration value can be added to the code named *hemoglobin concentration*) |
| Additional Characteristics | • Is able to assert other characteristics of a code that may be time- or context-dependent (e.g. new medical information may require updates to some codes) |
## Retrieval

| Analysis | Enables the consistent and reproducible storage of information, which is subsequently retrieved for analysis; this requires retrieval that allows the inclusion of subtypes and equivalent codes to be included. Equivalent codes may include:  
| - Codes represented in another (legacy) coding scheme  
| - Redundant codes that were inactivated  
| - Combinations of general codes and qualifying characteristics  
| - Analysis usually requires retrieval of selected records from a population of patient records; usually performed in batch |
| Patient Review | Enables the consistent and reproducible storage of information, which is subsequently retrieved for patient recall for preventive procedures or review; requirements similar to those for analysis |
| Decision Support | Enables the consistent and reproducible storage of information, which is subsequently retrieved for decision support  
| Requirements are broadly similar to those for analysis  
| - Decision support requires retrieval of selected records from an individual patient record  
| - Requires real-time processing to determine code meaning equivalence |
| Presentation | Enables the consistent and reproducible storage of information, which is subsequently retrieved for presentation  
| Requirements are similar to those for decision support  
| - Must be real-time, but usually involves filtering by broad categories of code; less precise than for decision support |
# Searches and Text Parsing

SNOMED CT facilitates searches for descriptions

- A simple keyword index may be generated from the descriptions and used for more effective searching although this may not be optimal due to:
  - Use of abbreviations
  - Word form variants
  - Word order variants
  - Word equivalences and combinations
  - Locally added mnemonics for frequently used descriptions
  - Composite coded meanings that can only be represented by:
    - Combinations of a code with one or more qualifying characteristics
    - Multiple codes related together by the patient record structure components
  - Searches with multiple redundant hits for a single code
    - When several synonyms of the same code match the search key
    - When techniques for word equivalences and combination are applied and return alternative descriptions related to the same code for two or more word equivalences
  - Searches with multiple redundant hits for a large number of closely related coded meanings
  - Search keys matching descriptions associated with a code with a more general meaning and many of its more specific hierarchical descendants

A further complication is the application of searches within subsets. This restricts the range of available concepts or codes; efficiency may depend on the relationships of keyword indices and subsets.

## Parsing or Encoding Free Text

- The use of natural language parsing to encode free-text derived from typing, scanning, or voice recognition is increasing; the text of descriptions and associated search indices may assist with this process.

# Implementation

## Terminology Services

- Terminology services should be implemented independent of application data; by individual applications or by terminology servers accessible by many applications.

## Advice

- Application data cannot be specified to the same level of detail as terminology services. It us dependent on the general functionality of the application and its record structure.
- Providing advice early in the SNOMED CT implementation process is required. This helps with some issues that may not be immediately apparent to developers.

## Limited Applications

- The advice provided should not place onerous requirements on applications with limited needs for the SNOMED CT terminology.
- It is inappropriate to have all-or-nothing requirements for SNOMED CT enabled applications.
## Legacy Data and Migration

<table>
<thead>
<tr>
<th>Code Recognition</th>
<th>• It should be possible to distinguish a code from an earlier coding schemes (SNOMED, Read Codes, or NHS Clinical Terms) from the identifiers used in SNOMED CT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Equivalence</td>
<td>• It must be possible to relate each code in early coding schemes (SNOMED, Read Codes, or NHS Clinical Terms) to a code in SNOMED CT</td>
</tr>
<tr>
<td>Query/Protocol Conversion</td>
<td>• There must be support to convert queries and protocols, based early coding schemes (SNOMED, Read Codes, or NHS Clinical Terms), to SNOMED CT compatible forms</td>
</tr>
<tr>
<td>Record Conversion</td>
<td>• It should be possible to convert legacy data, based on early coding schemes (SNOMED, Read Codes, or NHS Clinical Terms), to SNOMED CT compatible forms. This is subject to medico-legal constraints</td>
</tr>
<tr>
<td>Migration of Terminology-Dependent Products</td>
<td>• Projects in the UK NHS, that currently make use of Read Codes or NHS Clinical Terms, must plan migration to allow future use of SNOMED CT</td>
</tr>
</tbody>
</table>

## Data Structure

| Patient Record Architectures | • SNOMED CT is intended to represent clinical meanings in patient records  
|                             |   • A patient record consists of a series of related statements that are organized under headings  
|                             |   • The statements and headings may contain clinical codes derived from SNOMED CT  
|                             |   • Headings, and other contextual elements, may modify the meaning of related statements  
|                             |   • The relationship between a terminology, such as SNOMED CT, and a record architecture can be summarized as follows:  
|                             |   • SNOMED CT codes and terms may populate different elements in the record structure  
|                             |   • Different SNOMED CT codes may be applicable to different elements in the record  
|                             |   • Some codes may not be appropriate for inclusion in the record  
|                             |   • The meaning of a SNOMED CT code may be modified by its context within the record structure  
|                             | • SNOMED CT should be evaluated within the context of evolving standards for patient record architectures. Recommendations based on the evaluations may include:  
|                             |   • Possible changes to record architectures in order to realize benefits from SNOMED CT  
|                             |   • Changes to SNOMED CT to better fit into record structures  
|                             |   • Selecting SNOMED CT codes for use in specific record structure contexts |
Data Structure

Expression Coordination and Equivalence

• Some codes may be entered in a precoordinated or a post-coordinated manner

For example, "excision of ovary" might be entered by:

selecting the precoordinated code 83152002 |Oophorectomy (procedure)|,

or alternatively by selecting the codes for

71388002 |Procedure (procedure)| and adding the qualifying characteristics:

260686004 |Method (attribute)| = 129304002 |Excision - action (qualifier value)|
405813007 |Procedure site - Direct (attribute)| = 15497006 |Ovarian structure (body structure)|

• The coded meanings are stored in the forms entered. This may be using a single precoordinated code, a single post-coordinated expression, or a set of separate codes that together represent the clinical meaning.

• A retrieval query must therefore search for the precoordinated and all possible post-coordinated ways of expressing equivalent meanings. This can be done using the Expression Constraint Language (http://snomed.org/ecl) and a terminology service that can compute subsumption between expressions.

• These methods for retrieving records based on their clinical meaning rely on the formal definitions of SNOMED CT concepts being as complete as possible. Missing defining characteristics may result in problems with equivalence testing and therefore data retrieval.

Communication

Clinical Information

• The ability to communicate clinical information (represented by SNOMED CT) between applications must be supported

• Message specifications and other communication structures must accommodate SNOMED CT identifiers, and combinations of identifiers, in order to express postcoordinated coded meaning

Message Specifications

• Current message specifications (e.g. EDIFACT, HL7, and XML) use plain text files; SNOMED CT identifiers must use plain text so that they are appropriate for these messages

Postcoordinated Expressions

• Communication of postcoordinated expressions may be possible using specific qualifier fields in messages. This can also be accomplished by using syntactic representation of identifier combinations; these must be consistent with message syntax and field size limitations

Mapping

Classification

• Based on recorded codes, mapping tables are used to generate statistical and administrative data

• Automation of the process depends on the nature of the classification, the richness of the mapping table, and the functionality of the mapping software
## Mapping

| Grouping                                      | • Mapping tables are used to generate groupings for funding, administration, etc.
|                                             | • Mapping to a classification, then using the classification codes to generate groupings, is an alternative method |
| Communication Specifications                  | • Codes are mapped to specific values, in an enumerated list, associated with a message or communication specification
|                                             | • Recognizing these mappings may prevent double data entry, when sending or receiving such messages |
| Reference Works                               | • Codes are used to establish links with decisions-support protocols or other references
|                                             | • Mapping between these codes and reference sources may help to facilitate their use |

### Availability

| Limited Applications                          | • Applications vary in their ability to use terminological components
|                                             | • Special consideration may be necessary for applications that require only limited use of SNOMED CT |
| Concepts in Different Languages              | • Translating SNOMED CT into other languages is required
|                                             | • Multiple translations may support communication of clinical information across language barriers |
| Patients                                      | • Patients may be users of SNOMED CT if they record information in their own medical records
|                                             | • This may require limited licensing of SNOMED CT for populations, in general |

## Concept Model Overview

The **Concept Model** is used to specify logical definitions of **SNOMED CT** concepts. It is based on a combination of formal logic and editorial rules. It includes the attributes and values that may be applied to the concepts.

<table>
<thead>
<tr>
<th>Definition</th>
<th>Note</th>
</tr>
</thead>
<tbody>
<tr>
<td>The set of rules that determines the permitted sets of relationships between particular types of concepts</td>
<td>The <strong>Concept Model</strong> specifies the attributes that can be applied to concepts in particular domains and the ranges of permitted values for each of these attributes. There are also additional rules on the cardinality and grouping of particular types of relationships</td>
</tr>
</tbody>
</table>

## Root and Top-level Concepts

**Concept**

A **concept** is defined as a clinical idea to which a unique **concept identifier** has been assigned. Concepts are associated with **descriptions** that contain human-readable terms describing the concept.
Concepts are linked to their more general parent concepts directly above them in a hierarchy. More general meanings, are usually at the top of the hierarchy. Descending levels of the hierarchy contain more specific or specialized meanings.

Concepts are logically defined by their relationships to each other.

In SNOMED CT, the default meaning of a concept is defined above. However, a concept may have other meanings in SNOMED CT, such as:

- Abbreviated name for the concept identifier. For clarity, this is should be referred to as an identifier (ID), code, or concept identifier (ID).
- Idea or class of real-world entities (common usage meaning). For clarity, this is should be referred to as a clinical idea, clinical meaning, or code meaning.

Root Concept

The concept file includes a special concept referred to as the root concept. It is the single concept that is at the top of the SNOMED CT concept hierarchy. All other concepts are descended from this root concept via at least one series of relationships of the Relationship type 116680003 | Is a (attribute), i.e. all other concepts are regarded as subclasses of this concept. The root concept code is 138875005 | SNOMED CT Concept (SNOMED RT+CTV3). All other SNOMED CT concepts are subtypes of the root concept. Unlike other SNOMED CT concepts, the root concept is not a subtype of any other concept.
Top-level Concepts

Concepts that are directly related to the root concept by a single relationship of the Relationship type 116680003 |Is a (attribute)| are referred to as top-level concepts. All other concepts are descended from at least one top-level concept via at least one series of relationships of the Relationship type 116680003 |Is a (attribute)|, i.e. all other concepts represent subclasses of the meaning of at least one top-level concept.

Top-level metadata concepts

A concept that is directly related to the root metadata concept, 900000000000441003 |SNOMED CT Model Component (metadata)| by a single relationship of the relationship type IS_A. All metadata concepts are descended from at least one top-level metadata concept via at least one series of relationships with Relationship type IS_A. Metadata codes represent structural information about the terminology itself. The top-level metadata concepts represent broad groups of metadata.

Subtype relationships

Subtype relationships provide the main semantic hierarchy that relates concepts to one another. All active concepts, except the root concept, have subtype relationships with one or more concept. Each of these relationships indicate that a concept is a subtype of another concept.

Subtype relationships are expressed in the same way as other SNOMED CT relationships. They are identifiable by their fully specified names, e.g 116680003 |Is a (attribute)|.

For example,

- 53084003 |Bacterial pneumonia (disorder)| is a subtype of 233604007 |Pneumonia (disorder)| because it is a subtype of 312342009 |Infective pneumonia (disorder)| which is also a subtype of 233604007 |Pneumonia (disorder)|

Attributes

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Definition</th>
<th>Notes</th>
<th>Example</th>
</tr>
</thead>
<tbody>
<tr>
<td>Represents a characteristic of the meaning of a concept or the nature of a refinement</td>
<td>An attribute has a name which is represented by a concept. All of the concepts that can be used to name attributes are subtypes of the concept 410662002</td>
<td>Concept model attribute (attribute)</td>
<td>. An attribute is assigned a value (that creates an attribute-value pair) when used in the definition of a concept or in a postcoordinated expression. The permitted range of values for an attribute depends on the rules specified in the concept model.</td>
</tr>
</tbody>
</table>

Range

<table>
<thead>
<tr>
<th>Definition</th>
<th>Note</th>
<th>Example</th>
</tr>
</thead>
</table>
| A constrained set of values that the Concept Model permits to be applied to a specific attribute when that attribute is applied to a concept in a particular domain | The range of permitted values that can be applied to an attribute is typically defined to include concepts in one or more branches of the subtype hierarchy. The range for an attribute may include intensional or extensional definitions or both. An example of a range with an intensional definition is 370130000 |Property (attribute)| which has a range of <= 118598001 |Property (qualifier value)|. An example of a range with an extensional definition is 1148969005 |Has absorbability (attribute)| with range of 860574003 |Bioabsorbable (qualifier value)| OR 863965006 |Nonbioabsorbable (qualifier value)| OR 863968008 |Partially bioabsorbable (qualifier value)|. | 116676008 |Associated morphology (attribute)| is a subtype of 49755003 |Morphologically abnormal structure (morphologic abnormality)|.
Not all hierarchies in SNOMED CT have defining attributes. Many attributes apply to top-level domain hierarchies, some to more than one. Some attributes to a lower-level, or a more specific, domain hierarchy. Primitive concepts in some hierarchies may be attribute values in top-level hierarchies.

**Attribute definitions**
New attributes should include a text definition clearly indicating what the attribute means in the context of SNOMED CT.

**Attribute naming**
Attributes should be named to clearly communicate the property they specify and should refer to only one distinct property. The meaning of the attribute should not change if new values are added to the range.

If a new attribute is needed, look at existing unapproved SNOMED CT attributes and as well as other ontologies to see if a suitable attribute exists, including Basic Formal Ontology (BFO) ([https://basic-formal-ontology.org](https://basic-formal-ontology.org)), Relations Ontology (RO) ([http://www.obofoundry.org/ontology/ro.html](http://www.obofoundry.org/ontology/ro.html)), and Gene Ontology (GO) ([http://geneontology.org/](http://geneontology.org/)) as example ontologies to review.

Attributes should be named as verb senses, so that object-attribute-value relationships may actually be read. For example, a name of "Has filling (attribute)" is preferred over "Filling (attribute)" and "Has property (attribute)" is preferred over "Property (attribute)." Then a concept such as 464376000 |Saline-filled breast implant (physical object)| could be defined with the attribute "Has filling (attribute)" and a value of 387390002 |Sodium chloride (substance)|.

**Attribute datatype**
Many of the attributes in SNOMED CT have a range that includes SCTIDs as an allowed value. Attributes which have a binary (e.g., Boolean) value shall be valued using a descendant of 1119301001 |Boolean value (qualifier value)|: 31874001 |True (qualifier value)|; 64100000 |False (qualifier value)|. In the July 2021 release, a new attribute was created which uses Boolean style value: 1148965004 |Is sterile (attribute)|.

**Attribute hierarchy**
Selected SNOMED CT attributes have a hierarchical relationship to one another known as attribute hierarchies. In an attribute hierarchy, one general attribute is the parent of one or more specific subtypes of that attribute. Concepts defined using the more general attribute can inherit concepts modeled with the more specific subtypes of that attribute providing the attribute value is the same or a subtype of the attribute value used for the concept that is defined with the more general attribute.

Clinical finding and Event attribute hierarchies
- Associated with
  - Causative agent
  - Due to
  - Temporally related to
    - After
    - Before
    - During

Procedure attribute hierarchies
- Procedure Site
  - Procedure site - Direct
  - Procedure site - Indirect
- Procedure device
  - Direct device
  - Indirect device
  - Using device
    - Using access device
• Procedure morphology
  • Direct morphology
  • Indirect morphology

Body structure attribute hierarchy
• All or part of
  • Proper part of
    • Constitutional part of
    • Regional part of
      • Lateral half of
    • Systemic part of

Medicinal product attribute hierarchy
• Has ingredient (not used in the international edition)
  • Has active ingredient
  • Has precise active ingredient

Defining Characteristics

Role of defining characteristics
Defining characteristics represent the values of a range of relevant attributes. Depending on the nature of the concept, they may include etiology, topography, method, etc.

The attributes that can be applied depend on the domain of the concept. For example, a procedure may have a method, and a disorder may have an etiology, but a procedure cannot have an etiology, and disorder cannot have a method. Defining characteristics using a particular attribute will be applied consistently to all concepts to which it is relevant. Note that this design principle may not be fully realized for all attributes in each release.

Representation of defining characteristics
Defining characteristics are represented as relationships. The fields are used as follows:
• `SourceId` refers to the concept to which a defining characteristic applies;
• `TypeId` indicates the nature of the defining attribute;
• `DestinationId` refers to the concept that represents the value of that attribute.

Relationships
The defining characteristics can be divided into `116680003 | Is a (attribute) |` relationships and defining attribute relationships.

The IS_A relationship (also called supertype-subtype or parent-child relationship) builds the hierarchies in SNOMED CT. Every concept has at least one IS_A relationship to a supertype or parent concept.

**Exception**
138875005 [SNOMED CT Concept (SNOMED RT+CTV3)] has no supertype or parent relationship.

Each concept in SNOMED CT is logically defined through its relationships to other concepts. A relationship is defined as an association between a source concept and a destination concept. The type of association is indicated by an attribute concept. It is the relationships that make up the defining characteristics of the concepts. A defining characteristic is a relationship to a target concept that is always necessarily true for any instance of the source concept.

For example, the defining relationships of the concept 53442002 [Excision of stomach structure (procedure)] include:
• `116680003 | Is a (attribute) | = 65801008 | Excision (procedure)`
Qualifying Characteristics

A *qualifying characteristic* is expressed by an attribute-value pair. The attribute may have one value, from a range of values, based on the domain's concept model. If a particular qualifying characteristic is applied to a concept, the resulting expression represents a more tightly defined subtype of that concept.

Clinical expressions using SNOMED CT concepts can be of two types: precoordinated expressions, which use a single SNOMED CT concept identifier, and postcoordinated expressions, which contain more than one SNOMED CT concept identifier.

For example,

- It might be possible to qualify a disorder such as 53084003 | *Bacterial pneumonia (disorder)* | according to its clinical course (373933003 | *Acute onset (qualifier value)* | or 90734009 | *Chronic (qualifier value)*) or severity (255604002 | *Mild (qualifier value)*, 6736007 | *Moderate (severity modifier) (qualifier value)*, or 24484000 | *Severe (severity modifier) (qualifier value)*)
- 125605004 | *Fracture of bone (disorder)* can be refined by qualifying it with 12611008 | *Bone structure of tibia (body structure)* to represent the concept 31978002 | *Fracture of tibia (disorder)*

Authoring

Modeling philosophy of SNOMED CT

SNOMED CT authors use a zero-based, *proximal primitive* approach when modeling or editing logical definitions of concepts, i.e. a concept is newly defined, as opposed to inheriting the definition from the parent and then refining it. This is accomplished by assigning the immediate proximal primitive parent and attribute relationships based on their relevance to the defining characteristics of the concept, again, instead of relying on inheritance and refinement of relevant attributes from immediate, sufficiently defined supertypes.

The steps are as follows:

1. The author states the proximal primitive supertype/s.
2. The author states all of the defining *attribute-value pairs* required to express the meaning of the concept.
   a. An attribute-value pair is explicitly stated, even if it is already present on a supertype concept.
   b. The attribute-value pairs are grouped as required.
3. The classifier infers all appropriate proximal supertype/s.
   a. With sufficiently defined concepts, the subtypes are also inferred.

Advantages of the approach

- Enhances ability to maintain content
- Supports identification of equivalences

Content that does not conform

SNOMED CT contains content that does not conform to the current modeling patterns. A quality initiative is currently underway to correct these non-conforming concepts.

**Exceptions**

Exceptions exist where the current concept model is not expressive enough to represent critical defining characteristics of a concept that would allow for its sufficient definition.

For example, disorders where the clinical manifestations are variably present (i.e. genetic diseases)
Authoring information

Scope

International release criteria include affirmative answers to the following:

- Does it need to be understandable in electronic health applications in more than one national healthcare system?
- Can it be used in electronic health applications beyond a patient’s national healthcare system, i.e. if a patient were to travel or relocate to a different country?
- Is it useful in more than one national healthcare system?

The guiding principle underlying the creation of a clinical reference terminology is the facilitation of semantic interoperability. To this end, content in SNOMED CT must represent unambiguous, clinically relevant information which can be exchanged and understood internationally. A reproducible and consistent approach to incorporating terminology into electronic health applications is, therefore, mandatory.

The International Release includes content necessary for international conformance and interoperability (the International Release was formerly and is colloquially known as the core). The range of concepts, attributes, qualifiers, and other components of SNOMED CT is comprehensive compared to classification systems. This supports the terminological needs of those using SNOMED CT within electronic health applications.

Addition of new content to SNOMED CT requires careful consideration. Changes and additions to the International Release of SNOMED CT follow a formal process executed by SNOMED CT authors. For content to be included in the International Release, the following criteria must be met.

Broad use

It must be applicable within and across healthcare disciplines internationally.

Provision of use case

Changes and additions must follow SNOMED CT Content Request Service (CRS) Guidelines. It is very important to incorporate a clear justification for any change or addition request for the International Edition of SNOMED CT.

Principle of URU

- Understandable: The terminology must be able to communicate to recipients the intended meaning of the healthcare provider in terms that are unambiguous and comprehensible without reference to inaccessible, hidden, or private meanings.
- Reproducible: Concepts should be names that are human-understandable representations of the codes. It is not enough for an individual to say they think they understand a meaning. It must be shown that multiple people interpret and use the meaning in the same way. Can it be used in electronic health applications beyond a patient’s national healthcare system, i.e. if a patient were to travel or relocate to a different country?
- Useful: The meaning must have demonstrable use or applicability to health or healthcare.

Usefulness

Content submitted for inclusion in the International Release shall be required to pass a test for "usefulness." The usefulness test can be passed in more than one way. At least one of the following must be satisfied:

1. Content that is used by more than one major user (a National Release Center such as NHS, a vendor/supplier of clinical information systems with international scope, or a large intra-national system user such as VA or Kaiser) will be considered to have passed the "usefulness" criterion.
2. Data demonstrating significant frequency of use, or frequency of need, by a single user (single national center, or single vendor, or single health care system) can also be used as evidence in support of usefulness.
Additional means of passing the usefulness test may be added in the future. Submissions that pass the usefulness criterion must also pass understandability and reproducibility tests, and conform to style rules.

Naming of classes of things rather than instances
SNOMED CT concepts should name *classes of things*. Concepts that refer to a particular instance are unacceptable.

For example, *Doctor Jones pre-operative order set* should not be included because it is an individual instance, not a class.

References
Content must be submitted with:

- Definitions and literature references. All reference material must be publicly available. Wiki references are unacceptable.
- Evidence of international applicability. Without international applicability, a concept should, instead, be added to the submitter’s extension.

⚠️ Change Requests
For details on SNOMED International Content Request Service (CRS) Customer Guidance, see our [website](https://www.snomed.org) or the [Confluence site](https://confluence.snomed.org).

*(See also Appendix: Principles for Accepting Content in the International Release)*

Proprietary Names and Works
This section considers scope as it relates to the incorporation of proprietary names (e.g. brand names of drugs, devices, clinical forms or tools) into SNOMED CT.

Brand name of drugs and devices
Proprietary names are the names that have been assigned to products, usually drugs and devices, by their corporate producers. They do not require a license from the producer. It is both necessary and useful to include proprietary names in a health terminology. SNOMED International does not need to obtain the permission of the trademark owner simply to include a reference to the brand name drug in SNOMED CT. However, they should not be included in the International Release but instead in National Extensions. This is because proprietary names may refer to different products depending on the country and the meaning of these names are dependent on the country or jurisdiction in which the product is approved.

✅ Modeling
A brand or trade name may stand for a category of product and not the particular brand itself. These *proprietary* names may be included in the International Release as descriptions (non-FSN descriptions). They should not be included in FSNs.

For example,
- Kleenex, band aid, popsicle

Clinical forms, tools, or assessment scales
The owner of a form or tool may be an individual or organization that created it; the healthcare organization that employed the individual; or it may be a commercial organization to which the rights were assigned.
Names

Incorporating the name of a clinical form or tool (e.g. the XYZ Test (staging scale)), or the name of the score generated by a form or tool (e.g. the XYZ Test Score (observable entity)) into SNOMED CT does not require a license from the owner. Reference to the owner of a clinical form or tool loosely refers to the person or organization that owns the intellectual property rights of the form or tool. This may be the individual or group that originally created the form or tool, the organization that employed the creators, or a commercial organization to whom the creators assigned their intellectual property rights. It is possible that the owner holds a trademark (which may be registered or unregistered) representing the name or score, but simply incorporating that word into SNOMED CT does not infringe on the trademark.

Concepts representing questions, answers, or scores

A concept may be introduced into SNOMED CT that represents the text of questions, answers, or scores. For example, a form may include a question about a person's ability to dress and a range of possible answers. SNOMED CT may incorporate neither the text of the question nor any of the possible answers, but instead may incorporate a concept such as ability to dress. Similarly, if the form contains 20 questions, SNOMED CT may want introduce 20 concepts, for XYZTest_Result1, XYZTest_Result2, etc. to XYZTest_Result20.

The incorporation of a single concept into SNOMED CT, based on a question, answer, or score on a clinical form is highly unlikely to infringe on the copyright. However, if SNOMED CT systematically introduces a concept for every single question on a clinical form, it is likely to infringe on the copyright.

These concepts (e.g. ability to dress) may already exist in SNOMED CT, or they may be added in other contexts (This does not apply to concepts that represent specific questions within a form). This is unlikely to result in a copyright infringement.

Questions

A clinical form or tool, including the wording of the individual questions within the form or tool, is generally a literary work and qualifies for copyright protection (except in the case of the simplest of forms). The copying of all or any substantial part of a literary work, without a license from the owner, infringes on the copyright.

Answers

Certain questions may have a range of pre-determined answers. This could be as simple as Yes/No or a number within a specific range, but may also be more substantial (e.g. needs help cutting, spreading butter, etc., or requires modified diet). Incorporating very simple answers into SNOMED CT does not require permission. However, incorporating more substantial textual answers into SNOMED CT generally infringes on the copyright. This usually does not apply to individual answers, but it almost always applies when entire sets of answers are incorporated.

Scores

The principles that apply to individual answers also apply to the overall score generated by a clinical form or tool. The incorporation of numbers does not infringe on the copyright. However, when each possible score has an associated textual description and all possible scores and descriptions are incorporated into SNOMED CT, a license is required.

For example,

- 443807003 | EuroQol five dimension questionnaire (assessment scale) | is a SNOMED CT concept. However, these scores are subject to copyright protection, therefore cannot be added to SNOMED CT:
  - EuroQol Five Dimension (youth) doing usual activities score
  - EuroQol Five Dimension (youth) feeling worried, sad or unhappy score
Implementation of brand, trademark, and copyright

Names
The use of the name of a clinical form or tool or of a brand name drug will usually not infringe on the copyright. However, caution should be exercised by implementers who wish to use trademarks in a commercial context, i.e. a system that enables drugs to be purchased electronically. SNOMED International does not advise implementers on this matter, but recommends that implementers, who are in any doubt, contact the trademark owner.

In general, implementers should make no greater use of a trademark than is necessary. For example, displaying a graphical mark (e.g. a logo) on a screen or in printed material should be avoided.

Questions, answers, and scores
Implementers should manage questions, answers, and scores in the same way as SNOMED International (see above). When the incorporation of content from a clinical form or tool infringes on the copyright, the system the reproduces (by display or print) the content also usually infringes. This means that the license to incorporate content by SNOMED International should also cover the system implementer.

Preexisting terms
As noted above, terms in a clinical form may already exist within SNOMED CT, even though they have not been copied from the form. This is not copyright infringement by SNOMED International. If, however, a system implementer chooses to arrange a collection of these pre-existing terms in a way that reproduces all or a substantial part of a clinical form (e.g. by populating a drop-down box with all of the possible answers to a specific question on the form), that may infringe on the copyright.

Form structure
A system may reproduce the structure and layout of a clinical form on a screen display or printed output (e.g. to make the system more accessible to users who are familiar with a paper-based form). This may infringe on the copyright, unless the structure or layout is very minimal (e.g. a bulleted list). An implementer who wishes to emulate the look and feel of a clinical form should seek a license from the owner.

Algorithms or logic
System implementers may use the algorithm or logic inherent in a clinical form or tool (e.g. the method by which an overall score is calculated). For example, a clinical form may instruct the user to perform a mathematical operation on the individual answers to produce the overall score, and the same operation may be carried out by the system. The use of the algorithm or logic is an infringement. SNOMED International avoids such use and encourages system implementers to contact the owner to discuss possible infringement.

Adjudication for Content Requests
There are processes for making decisions about adding or changing content in SNOMED CT.

Change requests
All change requests, whether for new content or for change to existing content, go through a request submission approval process. It involves review by authors to determine that there is:

- International applicability
- Compliance with Understandable, Reproducible, Useful (URU) principles
- No duplication with existing content
- No link to existing larger projects, as detailed in a Content Tracker document
- No conflict with existing collaboration agreements (e.g. Logical Observation Identifiers Names and Codes (LOINC) agreement)
Legacy concepts

Legacy concepts, i.e. concepts not in the current draft/work-in-progress version of SNOMED CT, may not follow current guidelines. Requests based on legacy concepts are unacceptable.

Appeals, deferrals, and resolution

Appeals

Requests that are rejected may be appealed by the submitter.

Deferrals

Requests may be deferred for a number of reasons including questions about:

- How to model the concept; which attributes may be used
- Concept meaning
- Literature reference missing or inadequate
- Use case unclear
- Size of required change (attached to a Content Tracker)

Resolution

Resolution of deferrals may result in a decision delay requiring:

- A larger project or work item or
- Referral, internally, to other groups for decision. This depends on the complexity of the request and understanding of the wider impact.

Results

Results of adjudication are received by email from the Content Request System (CRS). Simpler issues can be resolved expeditiously (e.g. by a ruling from the Head of Terminology).

Descriptions

Descriptions

- A concept has multiple associated descriptions.
- Each description has a description type and a unique numeric description identifier.
- Fully specified name (FSN) and synonym (SYN) are description types in SNOMED CT.
- A preferred term (PT) is a synonym that has been marked as preferred in a particular dialect.
- Every concept may have only one preferred term in a specific dialect. Two preferred terms for the same language may coexist if they belong to two distinct dialects (e.g. variant US and GB of English language).

For example,

<table>
<thead>
<tr>
<th>Description type</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>FSN</td>
<td>Bleeding of mouth (disorder)</td>
</tr>
<tr>
<td>PT</td>
<td>Bleeding of mouth</td>
</tr>
<tr>
<td>SYN</td>
<td>Bleeding in mouth</td>
</tr>
<tr>
<td>SYN - US English</td>
<td>Oral hemorrhage</td>
</tr>
<tr>
<td>SYN - GB English</td>
<td>Oral haemorrhage</td>
</tr>
</tbody>
</table>
SNOMED CT Editorial Guide (2021-10-02)

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SYN

Stomorrhagia

241563001 | [Computed tomography of upper limb (procedure)]

<table>
<thead>
<tr>
<th>Description type</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>FSN</td>
<td>Computed tomography of upper limb (procedure)</td>
</tr>
<tr>
<td>PT</td>
<td>CT of upper limb</td>
</tr>
<tr>
<td>SYN</td>
<td>Computed tomography of upper limb</td>
</tr>
</tbody>
</table>

32849002 | [Esophageal structure (body structure)]

<table>
<thead>
<tr>
<th>Description type</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>FSN</td>
<td>Esophageal structure (body structure)</td>
</tr>
<tr>
<td>PT - US English</td>
<td>Esophageal structure</td>
</tr>
<tr>
<td>PT - GB English</td>
<td>Oesophageal structure</td>
</tr>
</tbody>
</table>

Fully Specified Name

Fully specified name (FSN) definition

A term unique among active descriptions in SNOMED CT that names the meaning of a concept code in a manner that is intended to be unambiguous and stable across multiple contexts.

⚠️ Precoordinated patterns

For information on acceptable precoordinated naming patterns, see [Pre-coordination Naming Patterns project](#). New content should conform with the naming patterns; however, legacy content may not.

An FSN is one type of description, unique among active descriptions in SNOMED CT. It provides the meaning of a concept so that it is unambiguous, stable across multiple contexts, and optimally understandable to those whose first language is not English. Consequently, it is not always clinician-friendly or in common use.

In the majority of cases, where the FSN is clinician-friendly and in common use, a description matching the FSN should be added to the concept. This description is not required to be the preferred term (PT). In certain instances, where the FSN does not provide a clinically useful description, a matching description without the semantic tag is unnecessary.

For example,

- FSN: Repair of common bile duct (procedure) - *the meaning*
- PT: Choledochoplasty - *commonly understood clinical name*

  Choledochoplasty is marked as *preferred* in the US English Language Reference Set; choledochoplasty is the *preferred term* for this *concept* in US English.

Each new content request should have an FSN that conforms to spelling, language, and style guidelines. It should also have SNOMED CT parent concepts that conform to editorial guidelines and show where in the hierarchy it belongs. In the Content Request System (CRS), if the meaning of the FSN is unclear or the parent codes are not provided, authors should request the information from the submitter.

A well formed FSN includes:

- Correct US spelling, not GB (General British) spelling
- Singular form, not plural form
- Procedures in present tense, not past tense
- A semantic tag in parentheses at the end

An FSN with an approved disjunctive (although not often used), e.g. Traumatic and/or non-traumatic injury of back (disorder), should have lower case and/or.
An FSN should not have:

- Abbreviations or acronyms
- Hyphens
- Duplicate concepts
- Ambiguity
- The word OR (not including the disjunctive and/or)
- Forward or backslash (/ \)
- Precoordinated numeric ranges
- Reference to a particular instance
- Reason or indication for a procedure, unless this directly impacts the method

Exceptions that should not be amended include:

- Trademark names
- Latin names of organism
- Scientific names

Structure, Structure of

When constructing the FSN for a disorder, finding, or procedure containing a body structure, the wording of the body structure should follow the naming convention of the body structure concept. However, it should not include the words structure or structure of.

For example,

- For the body structure concept, 266005 |Structure of lower lobe of right lung (body structure)|, a procedure with this body structure is 726425007 |Lobectomy of lower lobe of right lung (procedure)|.
- For the body structure concept, 74386004 |Nasal bone structure (body structure)|, a disorder concept with this body structure is 413878002 |Closed, displaced fracture of nasal bone (disorder)|.

Unique

The FSN is unique among active concepts. Creating a synonym to match the FSN is no longer mandatory because the SNOMED International Authoring Platform automatically creates a matching description to the FSN. Authors then determine the clinical usefulness of the matching description. Those that are useful are maintained in SNOMED CT; those that are not useful are removed. The Authoring Platform displays a warning when the matching description is removed; however, this does not prevent the author from saving the concept.

The FSN should provide a linguistic representation of the concept in an unambiguous way. It is considered an anchor for the representation of meaning of a concept, to which modelers can refer, when assigning a logic-based definition. The FSN does not necessarily follow the usual phrasing used in clinical practice; it may be phrased differently and may be longer and more fully spelled out in order to represent the meaning as clearly as possible and globally communicate the intended meaning of the concept.

The characters comprising the description, as well as case significance, must be taken into account to provide for a unique FSN. Uniqueness maintained through case sensitivity is handled by the "case significance indicator". It is possible to alter the semantics of concepts whose FSN uniqueness depends upon case significance.

For example,

38194003 |Weak e phenotype (finding)|
6800004 |Weak E phenotype (finding)|

The two referenced concepts above could easily be mistaken for duplicates if not for varying case sensitivity indicators that demarcate each concept's uniqueness.
Unambiguous

A single term may have more than one meaning. Therefore, FSNs should be checked for ambiguity.

For example, *immunosuppression* may mean the state of being immunosuppressed, or it may mean the application procedure of immunosuppressive therapy.

The following FSNs are clear and acceptable.

For example,
- Benign neoplasm of clavicle (disorder)
- Excision of cyst of spleen (procedure)

The following FSNs are ambiguous, and the concept should be inactivated.

For example,
- Standing in water side toward (finding); does not indicate *which side of what is toward what*
- Lumbar ache - renal (finding); does not convey whether the lumbar ache is specifically a renal etiology or is merely located in the renal area

Minor Changes - only the FSN changes but not the concept

Minor changes, those changes that do not change the meaning of the FSN, are allowed without inactivation of the concept. They may include:

- Capitalizing, i.e. from lower to upper case or upper to lower case
- Changing punctuation
- Changing spelling
- Replacing an acronym with its expansion (only if it is commonly understood and not ambiguous)
- Expanding an abbreviation
- Correcting word order without changing the meaning (only for an error)
- Correcting typos
- Removing articles, such as ‘the’, from concept string
- Aligning with editorial policy, e.g. changing *appendectomy* to *excision of appendix*
- Where a change to the FSN does not result in a change to the preferred term

Some FSN changes are necessary for style consistency; again, changes are only acceptable if the meaning does not change. They may include changing:

- Semantic tag type within a single top-level hierarchy
  For example,
  - A *finding* tag to a *disorder* tag
  - A *procedure* tag to a *regime/therapy* tag
- A substance or product name to reflect the International Nonproprietary Name (INN)
- The current scientific name of an organism (only applies to 410607006 |Organism (organism)| hierarchy)
Major Changes - When to inactivate the concept

Major changes to FSNs require inactivation of the concept. The following are examples of major changes, when:

- Changing the FSN changes the meaning
- FSN is ambiguous
- FSN meaning is more specific than the modeling; inactivation is determined case-by-case as this could simply be a primitive concept which cannot be defined
- Moving to a different top-level hierarchy
- Changing the common name to the scientific name
- Ancestors and descendants (if any) of the concept are inconsistent with what is implied by the FSN - inactivate concepts

International FSNs

The FSN for a concept in the International Release is designated an International FSN. The International FSN is considered the gold standard for interpretation of the meaning of the concept, from a linguistic standpoint.

The logical definitions, represented using the concept model, should represent the same meaning. Spelling of the International FSN follows United States (US) English spelling conventions. Other English language spelling and conventions, such as Great Britain (GB) English, may be represented in preferred terms and other descriptions. They should be appropriately tagged using the Language Reference Set mechanism.

For example,

- 191268006 Chronic anemia (disorder)
  - FSN: Chronic anemia (disorder)
  - US PT: Chronic anemia
  - GB PT: Chronic anaemia
- 414545008 Ischemic heart disease (disorder)
  - FSN: Ischemic heart disease (disorder)
  - US PT: Ischemic heart disease
  - GB PT: Ischaemic heart disease

Acronyms

Acronyms are easily misinterpreted. For this reason, all acronyms are unacceptable in FSNs.

For example, the FSN should be the expanded form, Computed tomography of chest (procedure), however as a preferred term, CT of chest (procedure) is acceptable.

If there is an acronym in an existing FSN, the FSN DescriptionId is inactivated and a new FSN is created (regardless of whether or not the acronym was in parentheses with the expanded form). The replacement FSN concept has the expanded description with the acronym entirely removed. Inactivating the ConceptId is not necessarily required, unless the FSN had significant ambiguity before changing it to its expanded form.
Imported FSNs

Before any changes are made to an FSN, imported directly with an extension (local) ID, the submitter should be notified and confirmation sought that no loss of meaning has occurred. This helps to ensure that the original meaning is understood and maintained. Authors should:

- Adhere to naming conventions.
- Advise the submitter of changes and confirm that they are acceptable.
- Check for existing concepts with the same FSN; the term may be added as a preferred term or synonym.

⚠️ **Original submitter**

Changes to existing SNOMED CT concepts do not necessitate notifying the original submitter.

Semantic Tag

*Semantic tags* are part of FSN descriptions. They are placed in parentheses at the end of FSNs when authoring concepts. They indicate the domain to which a concept belongs. For example, body structure, disorder, or specimen.

The purpose of semantic tags is to disambiguate concepts which have the same commonly used word or phrase. For example,

- Hematoma (morphologic abnormality)
- Hematoma (disorder)

The following table contains the semantic tags for each domain.

<table>
<thead>
<tr>
<th>Domain</th>
<th>Semantic tags</th>
</tr>
</thead>
<tbody>
<tr>
<td>Body structure (body structure)</td>
<td>• (body structure)</td>
</tr>
<tr>
<td></td>
<td>• (cell)</td>
</tr>
<tr>
<td></td>
<td>• (cell structure)</td>
</tr>
<tr>
<td></td>
<td>• (morphologic abnormality)</td>
</tr>
<tr>
<td>Clinical finding (finding)</td>
<td>• (finding)</td>
</tr>
<tr>
<td></td>
<td>• (disorder)</td>
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<tr>
<td>Environment or geographical location (environment / location)</td>
<td>• (environment)</td>
</tr>
<tr>
<td></td>
<td>• (geographic location)</td>
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<tr>
<td>Event (event)</td>
<td>• (event)</td>
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<tr>
<td>Observable entity (observable entity)</td>
<td>• (observable entity)</td>
</tr>
<tr>
<td>Organism (organism)</td>
<td>• (organism)</td>
</tr>
<tr>
<td>Pharmaceutical / biologic product (product)</td>
<td>• (clinical drug)</td>
</tr>
<tr>
<td></td>
<td>• (medicinal product)</td>
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<tr>
<td></td>
<td>• (medicinal product form)</td>
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<td></td>
<td>• (physical object)</td>
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<tr>
<td></td>
<td>• (product)</td>
</tr>
<tr>
<td>Physical force (physical force)</td>
<td>• (physical force)</td>
</tr>
<tr>
<td>Physical object (physical object)</td>
<td>• (physical object)</td>
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<tr>
<td></td>
<td>• (product)</td>
</tr>
<tr>
<td>Procedure (procedure)</td>
<td>• (procedure)</td>
</tr>
<tr>
<td></td>
<td>• (regime/therapy)</td>
</tr>
</tbody>
</table>
### Qualified value (qualifier value)

- (qualifier value)
  - (administration method)
  - (basic dose form)
  - (disposition)
  - (dose form)
  - (intended site)
  - (number)
  - (product name)
  - (release characteristic)
  - (role)
  - (state of matter)
  - (transformation)
  - (supplier)
  - (unit of presentation)

### Record artifact (record artifact)

- (record artifact)

### Situation with explicit context (situation)

- (situation)

### SNOMED CT Model Component (metadata)

- (attribute)
  - (core metadata concept)
  - (foundation metadata concept)
  - (link assertion)
  - (linkage concept)
  - (namespace concept)
  - (OWL metadata concept)

### Social context (social concept)

- (social concept)
  - (ethnic group)
  - (life style)
  - (occupation)
  - (person)
  - (racial group)
  - (religion/philosophy)

### Special concept (special concept)

- (inactive concept)
  - (navigational concept)

### Specimen (specimen)

- (specimen)

### Staging and scales (staging scales)

- (staging scale)
  - (assessment scale)
  - (tumor staging)

### Substance (substance)

- (substance)

### Preferred Term

A preferred term (PT) is the description that is deemed to be the most clinically appropriate way of expressing a concept in a clinical record. It represents a common word or phrase used by clinicians to name a concept in clinical practice or in the literature. It is the synonym that is preferred in a language or dialect.
The use of a description can vary between different languages, dialects and contexts. A description may be preferred in some dialects, acceptable in others, and may not be used in some dialects. A Language Reference Set is used to specify the descriptions that are preferred or acceptable in each language or dialect.

A concept may have two descriptions marked as PT, one for each language.

For example, 32849002 | Esophageal structure (body structure) | has
- PT: Esophageal structure (US)
- PT: Oesophageal structure (GB)

A PT for one concept may also be a synonym for another concept.

For example,
- 84162001 | Cold sensation quality (qualifier value) | has a preferred term of cold
- 82272006 | Common cold (disorder) | also has a synonym of cold

In both concepts, cold represents a common clinical phrase used to capture the meaning of the concept.

The PT is indicated by the acceptabilityId field, for a particular language or dialect.

**Synonym**

In SNOMED CT, a synonym (SYN) is a description that is an acceptable way to express the meaning of a concept in a particular language or dialect, i.e. it is a word or phrase, other than the FSN, that represents a concept. Unlike FSNs, synonyms are not required to be unique.

Each concept may have one or more synonyms.

For example,
- US English synonyms for 22298006 | Myocardial infarction (disorder) | are:
  - Myocardial infarction
  - Cardiac infarction
  - Heart attack
  - Infarction of heart
  - MI - myocardial infarction
  - Myocardial infarct

**Modeling**

A synonym may not change to, i.e. replace, an existing FSN.

**Duplicate terms as synonyms**

In most cases, it is unacceptable to add the same term as a synonym to more than one concept. However, some terms have more than one meaning and can be synonyms for more than one concept.

When concepts have the same term as synonyms, they are checked to determine whether or not they are duplicates. If they are duplicates, one concept is inactivated with a historical association link of SAME_AS to the other concept.

A synonym with a single meaning may be, erroneously, associated with more than one concept. If the concepts are not duplicates, the synonym should be retained with only one of the concepts and inactivated on the others.
Narrower synonym
When a synonym is more specific than the FSN, it does not have the same meaning, and should be inactivated. The description inactivation value of 723278000 |Not semantically equivalent component (foundation metadata concept)| is used.

For example:
- FSN: Removal of device (procedure)
- SYN: Replacement of prosthetic device (procedure) - more specific meaning than the FSN

Broader synonym
When a synonym is more general than the FSN, and there is no context in which it has the same meaning as the FSN, the synonym should be inactivated. The description inactivation value of 723278000 |Not semantically equivalent component (foundation metadata concept)| is used.

For example,
- FSN: Sprain (morphologic abnormality)
- SYN: Joint injury - more general meaning than the FSN

However, a more general synonym is acceptable when there is a context in which the synonym has the same meaning as the FSN.

For example:
- FSN: Entire fundus uteri (body structure)
- SYN: Fundus in the context of obstetrics - same meaning as the FSN

Definition
A definition is a textual description applied to some SNOMED CT concepts that provides additional information about the intended meaning or usage of the concept. Definitions are not mandated and are considered for addition on a case by case basis and if required to differentiate a concept from its related concepts.

Adding a definition to a concept provides additional clarity on its context of use. It "enhances" the definition provided by the modeled relationships whereby a term can be sufficiently defined logically, but the "words", which is how many look for and interpret meaning, may imply more or less specificity.

Definitions should be written as complete sentences beginning with a capital letter, ending with a period and marked CS, this is default for case sensitivity in DEF status.

For example,
- The definition for the concept 11530004 |Brittle diabetes mellitus (finding)| is:
  Frequent, clinically significant fluctuations in blood glucose levels both above and below levels expected to be achieved by available therapies.

  The definition should never be contradictory to the modeling.
The example, “raised blood pressure” is a commonly used phrase which in itself is ambiguous. Raised can mean "higher than a previous measurement"; "on the high side of normal range" or "above reference range". Because of that ambiguity, we may not be able to create a definition via the logical model, so would need a text definition to encourage a consistent use of the term.

Thus, if a term may be interpreted in multiple ways, but is intended to mean only one way in SNOMED CT, it needs a definition.

General Naming Conventions

In addition to the general naming conventions below, please also see any applicable naming conventions for specific hierarchies in their respective domain.

Generally, names should:

- Be consistent and reproducible
- Follow natural or human language when possible
- Be unambiguous to users
- Be clear for translation purposes

Naming conventions should not be based on word order preferences (e.g. to facilitate search or display). Creating multiple word order variants for these purposes is outside the scope of the International Release of SNOMED CT.

Articles

Descriptions should not include articles such as a, an, and the. There are legacy descriptions that contain articles such as the that will be corrected over time.

For example,

- Use description of Neoplasm of respiratory tract (disorder), not Neoplasm of the respiratory tract (disorder)
- Use description of Rupture of diaphragm (disorder), not Rupture of the diaphragm (disorder)

Abbreviations

Abbreviations are shortened forms of words or phrases, as such, they are open to interpretation. Consequently they are not permitted in fully specified names (FSN). They are not allowed in preferred terms (PT) or synonyms (SYN) unless they are accompanied by the fully expanded term. When no other terms are included in a description and the description begins with an abbreviation, then it is to be followed by a dash, and then the expanded word phrase.

For example,

- 273420000 | Disability assessment schedule (assessment scale) | has a synonym of DAS - Disability assessment schedule
Exceptions

Official names of organism, which is represented as organism preferred term, may include abbreviations. The abbreviations do not need to be accompanied by the fully expanded term.

- For example,
  - 448945001 [Campylobacter lari subspecies lari (organism)] has a synonym of Campylobacter lari subsp. Lari

Abbreviated organism part names are allowed in a preferred term (and other synonyms). The abbreviations do not need to be accompanied by the fully expanded term.

- For example,
  - 24771000087106 [Antigen of Streptococcus pneumoniae Danish serotype 1 capsular polysaccharide conjugated to Corynebacterium diphtheriae cross-reacting material 197 protein (substance)] has a synonym [Streptococcus pneumoniae Danish serotype 1 capsular polysaccharide antigen conjugated to Corynebacterium diphtheriae CRM197 protein] that includes CRM197 which is the abbreviated form for cross-reacting material.

The preferred term for allergen components in the Substance hierarchy follows the rules established by the World Health Organization/International Union of Immunological Societies Nomenclature Subcommittee: allergen names consist of the first three letters from the genus, one letter from the species epithet, followed by an Arabic numeral. e.g. 1157148000 [Arachis hypogaea 2 protein (substance)] has a preferred term of "Ara h 2".

Acronyms

An acronym is a specific type of abbreviation formed from the initial letters of words and is sometimes pronounced as a word (e.g. AIDS for Acquired Immunodeficiency Syndrome, NICU for Neonatal Intensive Care Unit). Acronyms have different meanings in different situations, therefore are open to interpretation and not approved for use in FSNs.

Fully specified names

Acronyms are not usually permitted in an FSN.

Exception: An acronym is allowed in an FSN when it has become a word in its own right, i.e. included in dictionaries; understood without expansion to its original full form.

For example,

- Concept 122456005 [Laser device (physical object)] uses the term "laser", which originated as an acronym for "light amplification by stimulated emission of radiation"

Preferred terms and synonyms

Acronyms are allowed in a Preferred Term or Synonym when followed by the expanded term. If the acronym stands alone (i.e. represents the entire meaning of the description without any other text), it is followed by a space, a hyphen, and another space, then the expanded term. The first word after the dash should be lower case as per usual capitalization rules.

For example,

- 30549001 [Removal of suture (procedure)] has a synonym of [ROS - removal of suture]
- 387727008 [Intermittent positive pressure breathing treatment (regime/therapy)] has synonym of [IPPB - intermittent positive pressure breathing therapy]

If the acronym forms only part of the description's meaning, it is followed by a space, then the expanded term in parentheses.

For example,
• |Nontraumatic AKI (acute kidney injury)| is a synonym for 140031000119103 |Acute nontraumatic kidney injury (disorder)|

Exceptions

The preferred term for imaging procedures involving imaging modalities commonly referred to by an acronym (such as CT, MRI, SPECT, PET) omits the expanded term after the acronym.

For example,

* |CT of head| is the preferred term for 303653007 |Computed tomography of head (procedure)|

Acronyms in rare and genetic diseases have been included for the concepts as part of the Orphanet project that do not include expansion of the abbreviation, as some of the diseases are commonly known by the acronym.

For example,

GLOW syndrome is a synonym of 782722002 |Global developmental delay, lung cysts, overgrowth, Wilms tumor syndrome (disorder)|. Although the FSN and other descriptions have full expansions, there is a description included for |GLOW syndrome|, which does not further delineate the GLOW acronym.

Second example,

BRESEK syndrome is a synonym of 717945001 |Brain anomaly, severe mental retardation, ectodermal dysplasia, skeletal deformity, ear anomaly, kidney dysplasia syndrome (disorder)|. Although the FSN and other descriptions have full expansions, there is a description included for |BRESEK syndrome|, which does not further delineate the BRESEK acronym.

Eponyms

Eponyms are names that are derived from proper names (usually the person who made the discovery or created the original description). They are found in many areas of medical terminology, including anatomic structures, morphologic abnormalities, diseases, findings, and procedures (e.g. Rutherford Morison’s pouch, vein of Galen, Aschoff body, Kell blood group, Down syndrome, Moro reflex, and Whipple procedure).

It is neither desirable nor possible to completely avoid using eponyms in a health terminology; although, if possible, they should be avoided. This helps to improve clarity of meaning and to facilitate translation to other languages. Fully specified names (FSN) should be full descriptions, whereas synonyms may be eponymous terms.

For example,

* Structure of great cerebral vein (body structure) has the synonym Vein of Galen
* Complete trisomy 21 syndrome (disorder) has the synonym Down syndrome
* Pancreatoduodenectomy (procedure) has the synonym Whipple procedure

It is permitted and encouraged to include eponyms as descriptions (non-FSN descriptions) whenever they are understandable, reproducible, and useful in a given context.

Exceptions

Exceptions require careful consideration since eponyms meanings may change over time. They are allowed when:

* The full description is exceptionally long and unwieldy (e.g. Hemi-Fontan operation (procedure) instead of bidirectional Glenn shunt with end-to-side anastomosis of proximal superior vena cava to right pulmonary artery with isolation from right atrium).
* The eponym is the only precise, clinically relevant name available.
* A non-eponymous name would necessarily be vague or subject to misinterpretation (e.g. Hodgkin lymphoma, Burkitt lymphoma).
**Preferred prepositions**

When constructing an FSN, the preposition 'of' is preferred over using the preposition 'in' to describe the morphology of an anatomic structure.

For example,

- *Cyst of scalp* should be used for the FSN and not *Cyst in scalp* because the latter may indicate a morphology within a layer of the structure, whereas 'of' indicates the morphology is within the region of the anatomic structure.

**Foundation hierarchies (body structure, substance, or organism) referenced in other hierarchy descriptions**

When creating the descriptions for a concept (e.g. a disorder concept) that name an entity such as a body structure, substance or organism, the conventions that are applied for naming the entity in the source hierarchy should be used.

For example,

- **11218009 | Infection caused by Pseudomonas aeruginosa (disorder) |
  
  This concept references the organism hierarchy in the causative agent of 52499004 | Pseudomonas aeruginosa (organism) | and uses that description in the FSN.**

- **143491000146106 | Felis catus protein (substance) |
  
  448169003 | Felis catus (organism) | has a preferred term of “Domestic cat”. As a result a related concept in the Substance hierarchy, 143491000146106 | Felis catus protein (substance) | has a preferred term of “Domestic cat protein”.**

- **337311000119101 | Blepharochalasis of left upper eyelid (disorder) |
  
  This concept references the body structure hierarchy in the finding site of | Structure of left upper eyelid (body structure) | and uses "left upper eyelid" in the FSN.**

- **12081931000119106 | Burn of left eye region (disorder) |
  
  This concept references the body structure hierarchy in the finding site of | Structure of left eye region (body structure) | and uses "left eye region" in the FSN.**

The descriptions should be context neutral for these foundation hierarchies. Where context is explicit for a disease or procedure, the preferred term from foundation hierarchies can be used instead. For example, procedure CT of abdomen indicates that the context of CT imaging is cross-sectional. The procedure site should be modeled with the cross-sectional abdomen. However, it is not necessary to change 'Computed tomography of abdomen (procedure)' to 'Computed tomography of cross-sectional abdomen (procedure)'. It is the same reason for the preferred term 'CT of abdomen'.

**Exceptions**

- Where an infection caused by a microorganism has a common name, the common name of the disease can be used in the preferred term if accompanied by the explicitly-stated organism.
• When defining concepts in other hierarchies by referring to an organism, it is not required to include the taxonomical rank of the organism in the FSN.
• There are instances where the requested term for a concept containing an organism common name doesn't correspond to specific taxa. Rather, the term is found as part of common names in multiple taxa belonging to a higher level taxon. For example, “parakeet” and “parrot” are not common names to any specific taxa, but are found as part of common names in multiple genera in 447329007 |Subfamily Psittacinae (organism)|. To avoid ambiguity, a grouper concept referring to high-level taxon should be created.
  • For example, In the Substance hierarchy, a grouper concept, 1149419004 |Psittacidae protein (substance)|, was created as a parent concept to the following two concepts:
    • 146711000146102 |Parakeet protein (substance)|
    • 146701000146104 |Parrot protein (substance)|

**Structure, Structure of**

Outside of the body structure hierarchy, concepts should not include the words structure or structure of in the concept descriptions.

For example,

• For the body structure concept, 266005 |Structure of lower lobe of right lung (body structure)|, a disorder concept with this body structure is 724056005 |Malignant neoplasm of lower lobe of right lung (disorder)|.
• For the body structure concept, 266005 |Structure of lower lobe of right lung (body structure)|, a procedure with this body structure is 726425007 |Lobectomy of lower lobe of right lung (procedure)|.

**Description Length Limitations**

In the rare event that the 255-character limit of the fully specified name is reached, standard naming conventions may be circumvented in order to adhere to the 255-term string limit. For example, the use of commas may be used instead of the word “and”.

**Case Significance**

Most SNOMED CT descriptions begin with an upper case letter in the SNOMED International Authoring Platform. Generally, the rest of the words in the description should be lower case except for abbreviations, proper nouns, i.e. names of people, organizations, taxonomic groups (e.g. species, genus, family), etc. The following values, as part of the SNOMED CT model component domain, provide details.

<table>
<thead>
<tr>
<th>Case Sensitivity</th>
<th>Values</th>
<th>Meaning</th>
<th>Examples</th>
</tr>
</thead>
</table>
| cl               | 900000000000020002 | Only initial character case insensitive (core metadata concept) | First letter of the description may or may not be capitalized while the case of the rest of the description cannot be changed | • Family history of Prader-Willi syndrome (situation)  
• Born in Australia (finding)  
• Neonatal jaundice with Dubin-Johnson syndrome (disorder)  
• Penicillin resistant Streptococcus pneumoniae (organism) |
### Case Sensitivity

<p>| | | |</p>
<table>
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<tr>
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<tbody>
<tr>
<td>CS</td>
<td>900000000000017005</td>
<td>Entire term case sensitive (core metadata concept)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Changing case may change the meaning of the term or is not commonly used</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• English as a second language (finding)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• pH measurement (procedure)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• mm (qualifier value)</td>
</tr>
<tr>
<td>ci</td>
<td>900000000000448009</td>
<td>Entire term case insensitive (core metadata concept)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Changing case does not change the meaning of the term</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Blood compatibility test (procedure)</td>
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<tr>
<td></td>
<td></td>
<td>• Bite of fish (event)</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Floor mat (physical object)</td>
</tr>
</tbody>
</table>

Special attention is to be paid to the possibility of altering the semantics of those concepts whose FSN uniqueness depends upon case significance.

For example,

The subtypes of 365638007 |Finding of Rh blood group (finding)| vary in meaning depending upon the description's case of the letters c, d, and e.

![Finding of Rh blood group (finding)](image)

**Children (10)**
- CDE haplotype (finding)
- CdE haplotype (finding)
- cDe haplotype (finding)
- cDE haplotype (finding)
- CDe haplotype (finding)
- cdE haplotype (finding)
- Cde haplotype (finding)
- cde haplotype (finding)
- Finding of Rh genotype (finding)
- Rh blood group phenotype (finding)
Figure 1: Stated view of 365638007 |Finding of Rh blood group (finding)| and subtypes

**Case sensitivity can be changed on an existing description without inactivating it.**

**The common name for |Structure of pharyngotympanic tube (body structure)| is eustachian tube. While the term originated eponymously, it is not the name of the person from whom the body structure was named after. Therefore, the case significance is ci for case insensitive.**

### Numeric values

Numeric values will not display differently if switched between upper and lower case, so numeric values should be treated as case insensitive characters in a term.

If a description begins with a numeric value and the word following the number does not begin with a capital letter, the case sensitivity indicator is ci for *Entire term case insensitive*.

For example,

- The concept 33635003 |Serotonin (substance)| has the synonym, 5-hydroxytryptamine. The description is recorded in SNOMED CT in lower case, not 5-Hydroxytryptamine, but the case sensitivity indicator is ci for *Entire term case insensitive*.

If a description begins with a numeric value and follows with an abbreviation that contains a capital letter, the case sensitivity indicator is cl for *Initial character case insensitive*.

For example,

- The concept 387407006 |Tioguanine (substance)| has the synonym, 6-TG. Apply the case sensitivity indicator of cl for *Only initial character case insensitive*. 
Special characters

Special characters such as <, %, >, ., &, ^, will not display differently if switched between upper and lower case, so numeric values should be treated as case insensitive characters in a term. The rules for numeric values apply similarly to special characters.

If a description begins with a special character and the word(s) and/or symbol(s) following the special character begins with a capital letter, the case sensitivity indicator is cl for Initial character case insensitive.

For example,

- The concept 277976001 | Less than 35 degrees C (qualifier value) has the synonym, <35 degrees C. The description starts with a special character that is case insensitive but contains an abbreviation "C" for Celsius that is case sensitive, so the case sensitivity indicator applied to the synonym is cl for Only initial character case insensitive.

Greek alphabet characters

Words derived from the Greek alphabet, for example, alpha, beta, delta, gamma, omega, and etc. are all case insensitive wherever they are in the description.

Assessment scales and staging systems

SNOMED CT descriptions representing assessment scales and staging systems should be capitalized per the name of the scale or staging system. Legacy concepts may not follow this pattern.

For example,

- Ages and Stages Questionnaires Third Edition (assessment scale)
- Fagerstrom test for nicotine dependence (assessment scale)
- National Cancer Institute histologic grading system (staging scale)
- Clark system for melanoma staging (staging scale)

Gram staining

Gram staining is a common laboratory technique used to differentiate bacteria based on their cell wall constituents. Laboratory test results may be Gram positive or Gram negative. The technique was developed by a Danish physician, Hans Christian Gram. Consequently Gram, when referring to the technique, should always begin with an upper case G.

Person Naming Conventions

Patient vs Subject

Descriptions should use the word subject, not patient, if required. Subject is broader than patient.

For example,

- 420058008 | Provider of history other than subject (person)

Subject refers to the subject of record, who may, in some circumstances, not be the patient.
Caregiver vs Carer

Descriptions with caregiver should be as follows:

- An FSN should use caregiver as (one word).
- There should be a synonym using carer.

For example,

- 425578005 |Caregiver able to cope (finding)|
- Synonym: Carer able to cope

Plurals

Fully specified names (FSNs)

In general, concepts are represented in the singular, rather than the plural.

For example:

- Disorder of lung (disorder), not disorder of lungs
- Acute cholecystitis due to biliary calculus (disorder), not biliary calculi

FSNs should not be plural unless the concept necessarily involves multiples.

Unintended plurals

Unintended plurals might be incorrectly interpreted. An unintended plural is the use of a plural when, in fact, there is only one entity.

Correct example,

- Multiple cranial nerve palsies; the word multiple indicates that there can never be just one, so the plural palsies is correct

Incorrect example,

- Trochlear lesion versus trochlear lesions; users would use this concept to refer to a single trochlear lesion, thus the plural form would be incorrect

Exceptions

Organizational nodes or grouper concepts may be plural.

For example,

- 234320004 |Procedures for splenic lesions (procedure)|
- 194732001 |Diseases of mitral and aortic valves (disorder)|, has IS A 195002007 |Multiple valve disease (disorder)|

A concept that necessarily involves multiples should have a plural FSN.

For example,

- Bilateral atrophy of testes (disorder)

⚠️ It is advisable to keep track of these exceptions in a separate subset or using a special term type, so that they can be excluded when the singular/plural distinction is important for mapping.

Punctuation and Symbols

Legacy content may not adhere to current guidelines and will be updated as resources allow.
Comma (,) A comma is allowed in an FSN when required for meaning or to add clarity.

For example,

- Computed tomography of head, neck, abdomen and pelvis (procedure)

A comma is not allowed to change sort order for use in the search function.

Unacceptable example,

- Frostbite, acute

Apostrophe (‘) Eponymous descriptions should not include an apostrophe or final s, unless the name normally ends in s. With rare exception, a concept with an eponym should have at least one description that follows this rule.

For example,

- Down syndrome, a synonym for Complete trisomy 21 syndrome (disorder)
- Sjogren syndrome (disorder)
- Meigs syndrome (disorder)

When common usage requires it, there should be at least one description that has the apostrophe s. For descriptions with a possessive apostrophe where the name normally ends in s, the apostrophe should follow the s.

For example,

- Alzheimer’s disease (disorder)
- Bowen’s disease (disorder)
- Meigs’ syndrome (disorder)

⚠️ Existing eponymous descriptions with the possessive s, but no apostrophe, need not be inactivated, but newly added descriptions should either have no s, or else include the apostrophe.

Special character (<, >, &, %, $, @, #) The special characters <, >, &, %, $, @, # are not permitted in FSNs. All instances of FSNs with these characters should be spelled out in full text.

For example,

- FD&C Yellow #2 should be FD and C Yellow Number Two

The characters &, %, and # are permitted in preferred terms or synonyms.

⚠️ The characters @ and $ are not used in any descriptions.

Hyphen and dash (–) A hyphen is used to join words and to separate syllables. Hyphens may be used in FSNs. There is no space either before or after a hyphen.

✔️ Hyphens should follow rules of style for the dialect and language in which the descriptions are used as found in such publications as the Chicago Manual of Style, the American Medical Association’s Manual of Style, a current medical dictionary, etc.

Unless used to prevent ambiguity, punctuation is to be used sparingly.

For example,
- Anti-infective agent (product)
- Zollinger-Ellison syndrome (disorder)
- Zellweger's-like syndrome (disorder)
- Tick-borne hemorrhagic fever (navigational concept)
- Phospho-2-dehydro-3-deoxygluconate aldolase (substance)
- Multidrug-resistant bacteria (organism)
- Pandrug-resistant bacteria (organism)
- Extended spectrum beta-lactamase-producing bacteria (organism)

A dash may be used to separate two phrases, to contrast values, or to show a relationship between two things. A dash should not be used in an FSN, with rare exception, because it may obscure the exact meaning of the description. The dash should be replaced with words that clarify the meaning. A dash is also used to separate an acronym from its expanded form when no other terms are included in a description.

For example,
- 273420000 Disablity assessment schedule (assessment scale) has a synonym of DAS - Disability assessment schedule
- 719977005 Communication Activities of Daily Living (assessment scale) has a synonym of CADL - Communication Activities of Daily Living

Exceptions

When there is a need to distinguish categories from more specific subtypes with the same name, a dash followed by the word category, may be used.

For example,
- 416500007 Malignant glioma - category (morphologic abnormality) distinguishes the category of malignant gliomas from those neoplasms that are called 74532006 Glioma, malignant (morphologic abnormality). The neoplasm called malignant glioma is one of several subtypes of 416500007 Malignant glioma - category (morphologic abnormality), and does not have the same meaning as the category itself.

Colon ( : )

In general, colons should not be used in fully specified names.

Exceptions

Colons are allowed in the FSNs of organisms, substances, or products where the colon is part of the name. They are also allowed in ratios and in tumor stages.

For example,
- Salmonella II 43: g,t: [1,5] (organism)
- Lidocaine hydrochloride 1.5%/epinephrine 1:200,000 injection solution vial (product)
- pT3: tumor invades adventitia (esophagus) (finding)

Colons may be allowed in non-FSN descriptions.

For example, to separate an abbreviation from the rest of a name or a specimen from the finding
- Urine: turbid (finding)

Forward slash (/)

The forward slash should not be used in FSNs. When the slash is part of the authoritative name (e.g. representation of heterozygosity in hemoglobinopathies), a hyphen (no space before or after) is used in the FSN. The forward slash, without spaces, may be used in a preferred term or synonym.
For example,
- FSN: Sickle cell-hemoglobin C disease (disorder)
- SYN: Hemoglobin S/C disease
- FSN: Per cubic millimeter (qualifier value)
- SYN: /mm³

Exceptions
A forward slash may be used to represent units of measure, official enzyme names, and laboratory test results. They may also be used in and/or when part of FSNs. There should be no space either before or after the slash.

For example,
- Nitroglycerin 0.3mg/hr disc (product)
- Ibuprofen 5%/Levomenthol 3% gel (product)
- Milligram/deciliter haptoglobin (qualifier value)
- Bone structure of head and/or neck (body structure)

A forward slash may be allowed in non-FSN descriptions in a variety of contexts. Some common examples of use are in acronyms with findings, and as an abbreviation meaning and/or concepts.

Plus sign ( + )
The plus sign is generally discouraged for use in descriptions, and legacy content still contains this symbol. However, some uses are allowed. Plus signs may be found in the product, disposition, and substance hierarchies.

For example,
- [H+/K+-exchanging ATPase inhibitor] is an acceptable synonym for 734582004 [Hydrogen/potassium adenosine triphosphatase enzyme system inhibitor (disposition)].

Caret symbol ( ^ )
A pair of caret symbols is used to enclose character strings that should display as superscript.

Current guidance for substance and product hierarchies is to not create new instances containing symbols for superscript and subscript.

The single caret is used to represent exponents, i.e. powers of, in alignment with the Unified Code for Units of Measure (UCUM) guidance on the use of powers of ten.

For example,
- 10^3 for the third power of ten

Pipe character ( | )
A description cannot contain a pipe character, |. Since the | is used to indicate the beginning and end of a description, it may cause confusion.

Umlaut ( ¨ )
An umlaut should only be accepted for terms that do not have equivalences in English. Synonyms without umlauts should be added to facilitate searching in English.

For example,
- 83901003 [Sjögren's syndrome (disorder)] and one of its synonyms, Sjogrens syndrome
Sentence Types

Concepts should be names or short noun phrases. Full statements or sentences are unacceptable.

Procedure concepts should not contain phrases that can be categorized as a sentence function type, i.e. imperative, declarative, interrogative, or exclamatory. A procedure description should be a noun phrase that names the procedure, and should not contain information that it was done, or is to be ordered, carried out, or planned.

For example,

- 11227005 | Excision of ganglion of tendon sheath of hand (procedure) | is a noun phrase giving the proper description for the procedure

Unacceptable example,

- Hand tendon ganglion excised (situation) indicates the procedure was done, as a past tense declarative statement

  This is a situation with explicit context, not a procedure.

US vs. GB English

All fully specified names (FSN) should be represented in US English. When there is a difference between the US and GB spelling, there should be US and General British (GB) preferred terms (PT) and/or synonyms (SYN).

For example:

- FSN: Benign tumor of endocrine pancreas (disorder)
- PT-US: Benign tumor of endocrine pancreas
- PT-GB: Benign tumour of endocrine pancreas

Proper nouns

Where an FSN represents the name of an organization or trademarked name, a synonym with variant GB or US spelling is not required.

---

**References for Spelling**

<table>
<thead>
<tr>
<th>References</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>US-GB differences</strong></td>
<td>Wikipedia, the free encyclopedia*&lt;br&gt;*Note: Wikipedia may be used as a starting point, or source, for authoritative references, but not as an actual reference</td>
</tr>
<tr>
<td><strong>US Medical English</strong></td>
<td>Stedman’s Medical Dictionary&lt;br&gt;Merriam-Webster Online Dictionary&lt;br&gt;American Medical Association (AMA) Manual of Style</td>
</tr>
<tr>
<td><strong>GB English</strong></td>
<td>Dorland’s Medical Dictionary - medical terminology&lt;br&gt;Chambers 21st Century Dictionary - general</td>
</tr>
</tbody>
</table>
Principles for selecting preferred spelling variants

SNOMED CT may include (or add) more than one description, each with a different spelling for a given concept. That is, if the above references provide evidence of acceptability in the dialects for which they are being added.

For spelling of preferred terms in a dialect, where the reference sources provide multiple options, a judgment about the most common spelling may be needed. This may be determined by reviewing journal articles containing the word in question.

- Articles should be from highly cited journals, e.g. BMJ (for British English) or NEJM or JAMA (for US English).
- For concepts that are not clinical, appropriate scientific journals should be consulted, e.g. Science (US publisher) or Nature (UK publisher).

Fetal vs. Foetal

Fetal is the preferred term in both the US and GB language reference sets. Fetal is acceptable in GB synonyms. Foetal is not acceptable for US language but acceptable for GB language.

Action Verbs

Action verbs should be written in noun form within SNOMED CT descriptions. This most often means the verb will end with a suffix of –tion, -sion, -ment, -al, -ence, or -ance.

For example,

- Destruction instead of destroy
- Incision instead of incise
- Replacement instead of replace
- Removal instead of remove
- Maintenance instead of maintain

However, the root form of the verb may be used when it does not make a word when ending in noun suffixes.

For example,

- Control
- Release
- Care

Lastly, the verb with a suffix of –ing may be used when the root form of the verb may cause ambiguity in the meaning, i.e. the root form of the verb could also be a physical object.

For example,

- Wiring instead of wire
- Suturing instead of suture
- Splinting instead of splint
- Mapping instead of map
- Grafting instead of graft

Exceptions,
Common usage may dictate some exceptions.

For example,

- **Repair** instead of repairment

  Although ‘repairment’ may be considered a valid word, its use has fallen out of common usage in comparison to ‘repair’.

Check for approved and unapproved naming patterns in the [Precoordination Naming Pattern Project](#).

### Past tense

A past tense verbal phrase should not be used to name a procedure, since it indicates that the procedure was done in the past.

Unacceptable example,

- *Hand tendon ganglion excised* indicates the procedure was done, as a past tense declarative statement.

However, the following is an acceptable example using a noun phrase,

- 11227005 | Excision of ganglion of tendon sheath of hand (procedure)

**Situation hierarchy**

Existing descriptions containing past tense verbs should be moved to the 243796009 | Situation with explicit context (situation) hierarchy.

### Numbers and Numeric Ranges

#### Roman numerals versus Arabic numbers

Use the most common representation found in literature for the fully specified name. Use the alternative representation as a synonym, if it is also represented in the literature. If neither representation is common, use the Arabic representation.

For example, in the *AMA Manual of Style*, cancer stages are expressed with the use of capital Roman numerals: stage I, stage II, stage III, stage IV. The term, "stage 0", usually indicates carcinoma in situ. Histologic grades are expressed with Arabic numerals, e.g., grade 2.

#### Numeric ranges

In general, content that depends on numeric ranges should not be used for precoordination.

For example,

- There may be too many possibilities
  - A finding of number of lesions might have ranges of 1, 2 to 5, and greater than 5; 1 to 2, 3 to 10, and greater than 10, or etc.

- There may be possible changes to reference ranges or systems of units
  - The normal serum sodium concentration is usually defined as 135 to 145 mEq/L. Low serum sodium should not use the phrase *serum sodium less than 135 mEq/L*. (It should use a phrase such as *serum sodium concentration below reference range*)
  - A body mass index (BMI) score as an indicator of obesity
SNOMED CT Editorial Guide
(2021-10-02)

General Modeling

SNOMED CT is arranged as a polyhierarchy. A hierarchy is defined as an ordered organization of concept codes linked together through IS A relationships. Concept codes are linked to their more general parent concept codes directly above them in a hierarchy. Concepts with more general meanings are usually located at the top of the hierarchy and then at each level down the hierarchy the meanings become increasingly more specialized.

For general modeling information, use the following links to jump to the following pages:

Changes to Components

Concepts, descriptions, and target values may be changed for a variety of reasons.

Concept Inactivation

Concept inactivation values

Depending on the reason for inactivation, a specific Inactivation reason has to be selected.

<table>
<thead>
<tr>
<th>Inactivation reason</th>
<th>Association type</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ambiguous</td>
<td>Possibly equivalent to</td>
<td>• The concept has been made inactive because it is inherently ambiguous. This may be because of an incomplete fully specified name or because it has several associated terms that are not regarded as synonymous or partially synonymous.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• The possibly equivalent target is an active concept that represents one of the possible meanings of the inactive concept.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Multiple rows may be used to refer to each of the possible replacement targets for the ambiguous concept.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Previously referred to as May Be A.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Ambiguous concepts with a single target</td>
</tr>
</tbody>
</table>

A single target may be justified in the following situations:

• If one of the meanings of the ambiguous concept is not clinically useful.
• The FSN represents a classification concept, such as ‘other’, ‘NOS’.

Exception: acceptable numeric range

A standard definition with a fixed numeric range, i.e. the range is an explanation or definition of the score, may be acceptable.

For example,

• A histologic scoring system with a score of 1 when there are 0 to 5 mitoses per high power field, and a score of 2 when there are 6 to 10, and etc
• The Tumor, Node, Metastases (TNM) Classification of Malignant Tumor.

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### Inactivation of duplicate concepts

**Prior to inactivation**
- Check to see if the two concepts are true duplicates, i.e. semantically equivalent.
- Decide if the semantic meaning of the two concepts is the same.
- Review ancestors and descendants (if any) of the concept. Are they inconsistent with what is implied by the FSN? If so, inactivate the concept.

**Inactivation**
- Keep the more specific FSN and keep the concept ID.
  - Note: Implementers do not see the modeling. Hence there should be more weight in the meaning of the FSN, rather than the underlying modeling.
  - If appropriate, add the inactivated FSN as a synonym for retained concept.
  - Add the synonyms from the inactivated concept, where they are semantically equivalent,

**Consider**
- Inactivating the concept with fewer subtypes. This will simplify the process and minimize the amount of rework required.
- If needed, the retained FSN should be reworded to align with current policy. If required, modeling should also be corrected.

**Inactivated concept**
- Add the inactivated descriptions as synonyms (if the meaning is the same) to the retained concept.
- The inactivated concept should be marked as ambiguous, if it has an unclear meaning.
Inactivating classification-derived terms

SNOMED CT has many concepts derived from classifications that describe a clinical condition with an added exclusion, such as 90768003 |Contusion of brain without open intracranial wound (disorder)|. These “without x” terms are not clinically useful. When inactivating these concepts, use Ambiguous as the inactivation reason with the association type of Possibly equivalent to the parent concept, which is the clinical condition without any context.

For example,

When inactivating 156322003 |Pilonidal sinus without abscess (disorder)|, use the inactivation reason Ambiguous with Possibly equivalent to association type pointing to |Pilonidal sinus|.

Description Inactivation

**Description inactivation values**

Depending upon the combination of the type of component and the reason for inactivation, a specific Inactivation reason has to be selected.

<table>
<thead>
<tr>
<th>Inactivation value</th>
<th>Definition</th>
<th>Example</th>
</tr>
</thead>
<tbody>
<tr>
<td>Not semantically equivalent component (foundation metadata concept)</td>
<td>A description does not represent the same meaning as the concept’s Fully Specified Name (FSN)</td>
<td>Removal of device (procedure) has a synonym, Replacement of prosthetic device (procedure), which should be inactivated because the synonym has a more specific meaning than the FSN.</td>
</tr>
<tr>
<td>Outdated component (foundation metadata concept)</td>
<td>A component is no longer current, useful, appropriate or acceptable</td>
<td>The synonym Compression facies was inactivated from the concept’s more modern description of Facial asymmetry.</td>
</tr>
<tr>
<td>Erroneous component (foundation metadata concept)</td>
<td>A component contains a technical error</td>
<td>Case significance error: Alpha should have a lower case a&lt;br&gt;Spelling error: Asthma misspelled as Assthma</td>
</tr>
<tr>
<td>Nonconformance to editorial policy component (foundation metadata concept)</td>
<td>A component fails to comply with the current editorial guidance</td>
<td>The concept Urine: turbid (finding) was inactivated and replaced by 167238004</td>
</tr>
</tbody>
</table>

Order of selection of inactivation values

When there is more than one reason to inactivate a description, the order of preference for the inactivation value is as follows:

1. 723278000 | Not semantically equivalent component (foundation metadata concept)|
2. 9000000000000483008 | Outdated component (foundation metadata concept)|
3. 9000000000000485001 | Erroneous component (foundation metadata concept)|
4. 723277005 | Nonconformance to editorial policy component (foundation metadata concept)|
Corresponding association requirements

Only the description inactivation value of *Not semantically equivalent* requires an association type; the association type is *Refers to* and necessitates the reference to an active SNOMED CT concept. The other three description inactivation values (outdated, erroneous, nonconformance) do not require an associated concept.

Target Values

Considerations for **range** Concepts

Concepts that are used as a target value in an attribute relationship impact the placement of the source concept of the relationship. Some concepts, for example, those in the Qualifier value hierarchy, are created to support the definition of other concepts.

**Review after addition of new attribute value**

Creation of a new concept that will be used as the target value in an attribute relationship requires an author to determine if there are active concepts in the *domain* hierarchy that should also use the new concept as a target value.

For example,

```
The creation of a concept 713295009 | Surgical replacement - action (qualifier value) | would require a review of current active concepts that represent surgical replacement procedures; that were previously modeled with the attribute relationship Method = Replacement - action.
```

```
A concept that represents a surgical replacement procedure that currently has the relationship Method = 282089006 | Replacement - action (qualifier value) |, would require inactivation of the relationship and creation of a new relationship Method = 713295009 | Surgical replacement - action (qualifier value). |
```

Conjunction and Disjunction

In *SNOMED CT*, *and* is used in descriptions to represent the operator for logical conjunction. Concepts with the disjunctives *(or, and/or)* are unacceptable. Instead, there should be separate concepts. There are limited exceptions where *and/or* is used to represent the operator for inclusive disjunction. This helps to avoid confusion with the literal use of *or* in common language, i.e. only one of two operands is true; rarely both operands are true.

<table>
<thead>
<tr>
<th>Conjunction and Disjunction</th>
<th>and</th>
<th>or</th>
<th>and/or</th>
</tr>
</thead>
<tbody>
<tr>
<td>SNOMED CT</td>
<td>Conjunction: And</td>
<td>Exclusive disjunction: Or</td>
<td>Inclusive disjunction: And/or</td>
</tr>
<tr>
<td></td>
<td><em>A set of operands is true, if and, only if all of its operands are true</em></td>
<td><em>Either A or B is true but not both</em></td>
<td><em>A set of operands is true, if and, only if one or more of its operands is true</em></td>
</tr>
<tr>
<td></td>
<td><em>A and B are true</em></td>
<td></td>
<td><em>Either A or B is true</em></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td><em>or</em></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td><em>Both A and B are true</em></td>
</tr>
</tbody>
</table>

Disjunctives

Concepts with disjunctives *(or, and/or)* in disorders and procedures often involve one or more body structures.

For example,

```
65966004 | Fracture of forearm (disorder) |
```

The concept does not specify which bone of forearm is fractured. It is a break in *one or both of the radius and/or ulna* per the ICD definition. It would subsume fracture of radius, fracture of ulna, and fracture of both radius and ulna.
Exclusive disjunction ("or" only) is used when either operands is true but both cannot be true.

For example,

- 417163006 | Traumatic or non-traumatic injury (disorder)

Disjunctives are unacceptable with limited exceptions below. Instead of disjunctives, there should be separate concepts when possible.

### Exceptions

Disjunctives may be used if the:

- The referent is a single thing, but there isn't a name for it.
  
  For example,
  
  - 774007 | Structure of head and/or neck (body structure)

- The concept is an intensional navigational aggregate.
  
  For example,
  
  - 707861009 | Structure of skin and/or skin-associated mucous membrane (body structure)
  - 768845000 | Xanthine and/or xanthine derivative (substance)
  - 767271006 | Lead and/or lead compound (substance)

- The concept is based on an authoritative source but not a classification system.

### Modeling

The use of and/or in a description with disjunction should be lower case.

Anatomical structure hierarchy

Conjunction and disjunction are commonly used in the anatomical structure hierarchy.

For example, 419605007 | Structure of ankle and/or foot (body structure) represents adjacent regions of ankle and foot by a single concept. It is inclusive disjunction because any structures of ankle, foot, or both are true subconcepts. Entire ankle and foot as a conjunction means the ankle and foot as a whole. The concept represents the entirety of this single region though there is no dedicated name.

Following the anatomy SEP (Structure/Entire/Part) model, structure means all or any part of an anatomic entity, which is inclusive disjunction. Structure of ankle and foot represents all or part of entire ankle and foot. Therefore, any structures of ankle, foot, or both are true subconcepts of structure of ankle and foot. Structure of ankle and foot has the same meaning as structure of ankle and/or foot. The use of and/or is actually redundant for structure concepts like 419605007 | Structure of ankle and/or foot (body structure).

### Inclusive disjunction

Structure of ankle and foot was previously used. These descriptions were changed to and/or to explicitly indicate inclusive disjunction. This supports users who are unfamiliar with the interpretation of structure in the SEP model.

**And**

The and represents conjunction in disorders and procedures that can be interpreted as co-occurrent. It can be read as both in common usage. It would be all if it refers to more than two disorders or procedures.

For example,

- 75857000 | Fracture of radius AND ulna (disorder) represents the occurrence of a fracture of radius and a fracture of ulna at the same time or event. In other words, fracture of both radius and ulna. The concept should be modeled using two finding site relationship groups: Bone structure of radius in one and Bone structure of ulna in the other.
General Concept Inclusions - GCIs

Draft guidance

See the background, use cases, and examples for general concept inclusion axioms as well as explanation of the definition status at General Concept Inclusion 0.01.

Authoring Platform User Guide for GCIs

Reference the SNOMED International Authoring Platform User Guide for technical information describing how to add an additional axiom and general concept inclusion.

GCI display in the browser

A concept with GCIs will display in the browser in the stated view only.

For example,

Below is 417163006 |Traumatic or non-traumatic injury (disorder) in the stated view with the GCIs appearing to the right of the concept:

![Diagram of stated view with GCIs](image)

Here is the same concept in the inferred view without the GCIs appearing:

![Diagram of inferred view without GCIs](image)
Modeling concepts with a GCI-modeled supertype

General concept inclusions allow multiple definitions of a concept. A group of subtypes may be defined using GCIs and be considered subtypes of the parent concept without fully defining that parent concept. That parent concept could have multiple definitions, each of which is valid but none of which completely describes the parent concept on its own.

When modeling a concept that will be inherited by a GCI-modeled concept, there is no need to add the GCI-modeled concept as a stated parent, even if that GCI concept is primitive, if subsumption still occurs given the GCI axiom.

For example,

- Thunderstorm asthma (disorder) below shows modeling stating two primitive parents. Allergic condition (finding) is modeled with a GCI as notified by the salmon pink color. The left side of the diagram shows the inferred view.

Because the GCI-modeled primitive parent is unnecessary to state in the model, the diagram below shows modeling of the concept with the absence of Allergic condition (finding) as a parent, and yet the inferred view diagram on the left is still the same as compared to the incorrectly modeled diagram above.
Alternatively, if a GCI-modeled parent will not subsume an appropriate child concept, then the GCI-modeled concept should be stated as a primitive supertype.

For example,

28078000 [Closed fracture of shaft of bone of forearm (disorder)] must have the GCI-modeled concept stated as a primitive supertype, or else the concept would not be subsumed by [Fracture of shaft of radius and/or ulna (disorder)].
Defining GCI-modeled concepts

Though most are primitive, it is possible to define concepts modeled with GCIs. A concept must continue to meet the necessary conditions in order to be considered defined. GCIs can be added to extend the subtypes a defined concept will infer when appropriate.

For example,

Peripheral arterial occlusive disease (disorder) is defined as an obstruction of a peripheral artery. In addition, there are three general concept inclusions.
Points to Consider

- GCIs are not restricted to particular hierarchies; they can be used as applicable in any hierarchy that has a concept model.
- The Authoring Platform does not currently have the ability to create templates that include GCIs.

Grouper Concept

For hierarchies with a concept model, the usefulness of fully-defined groupers is limited to convenience groupings based on particular use cases. They may be added if they provide demonstrable benefit to organizing and navigating the terminology.

Grouper concepts provide a definition for subtypes that are always and necessarily true. The grouper concept must be sufficiently defined and clinically useful for the purpose of organizing content for an intensional reference set (e.g. disease of colon and all of its descendants) or in Expression Constraint Language (ECL), 128524007 | Disorder of colon (disorder)|
Navigational concepts

Grouper concepts should not be confused with navigational concepts. Navigational concepts were created to group other concepts without explicit regard for defining attributes (since there were none). Their purpose was to provide top level groupers for subsets and reference sets used in implementations. Because the Reference Set mechanism is now available, there is no longer a need for navigational concepts in the International Release; however, they can be added at the national or lower level.

In the past, there was an indiscriminate move of concepts in and out of the navigational concept hierarchy based arbitrarily on use cases by those users organizing concepts based on a particular classification that was wanted. The navigational concept hierarchy was useful to group things into a particular domain. The problem is that many of these are domain-specific and cannot be generalized. For example, mosquito-borne diseases will vary depending on the location of the user. It is difficult to classify the complete instance of these as well. Potential children would have to be manually assigned.

Because this is a primitive hierarchy and subtypes will not auto classify, much work would be required to reorganize hierarchies and maintain the use of navigational concepts. Inactivating concepts may be met with requests to create intermediate primitives. The Content Managers Advisory Group [CMAG] at Use of navigational concepts is being consulted regarding current use of navigational concepts.

As 363743006 |Navigational concept (navigational concept)| is within the 370115009 |Special concept (special concept)| subhierarchy, please see that section of the Editorial Guide at Special Concept.

Intermediate Primitive Groupers

Intermediate primitive groupers add a substantial management burden, thus, are discouraged. They may however be added on a case-by-case basis with approval from the Head of Terminology or the Principal Terminologist when for example:

- The Concept model is not robust enough to support the full definition of a subset of terms e.g. genomics (i.e. genetic diseases for which we cannot state, the majority of cases of this disease present with X)
- There are variances in the clinical manifestations

If an existing intermediate primitive concept cannot be sufficiently defined and has only one subtype, is not used to model another concept nor demonstrably clinically useful, it should be inactivated.

Rules for grouper concepts

A grouper concept that is added to SNOMED CT must adhere to the following rules:

- The concept must not be created with the hierarchical tag, (navigational concept).
- The concept must use the semantic tag for the relevant hierarchy e.g. (finding), (procedure).
- The concept must not have stated subtypes. All subtypes must be inferred by the classifier.
- The grouper concept will ONLY be added if it can be sufficiently defined.

Where grouper concepts already exist, the following criteria apply:

- If it can be sufficiently defined, remodel it and reassign existing stated subtypes to a new proximal primitive parent.
- Identify primitive concepts that cannot be sufficiently defined for additional review.

Modeling

If the addition of a grouper concept duplicates a concept in the 363743006 |Navigational concept (navigational concept)| hierarchy, the navigational concept should be inactivated.
Sufficiently Defined vs Primitive Concept

Sufficiently defined

A concept is sufficiently defined if its defining characteristics are adequate to define it relative to its immediate supertypes. A sufficiently defined concept is defined in the context of its hierarchy. See main glossary entry for sufficient definition.

Primitive

A concept which is not sufficiently defined is primitive. A primitive concept is a formal logic definition that is inadequate to distinguish it from similar concepts. A primitive concept does not have enough defining relationships to computably distinguish it from more general concepts (supertypes).

Proximal Primitive Modeling

See glossary for definition here: proximal primitive (PP)

- For some, but not all concepts, it is a top level concept e.g. Procedure.
- The proximal primitive supertype may also be an intermediate primitive concept located between the top level concept and the concept in question.
- There may be more than one proximal primitive supertype for a concept.

The approved modelling approach is to use:

- Proximal primitive supertypes
- Attribute-value pairs sufficient to define the meaning
  - An attribute-value pair is explicitly stated for the concept, even if it is already present for a supertype concept.
  - Attribute-value pairs are grouped as required.

The classifier infers all appropriate proximal supertypes. With sufficiently defined concepts the subtypes are also inferred.

For example,

- The stated view of 702499000 | Computed tomography of humerus (procedure). The PP supertype for this concept is 71388002 | Procedure (procedure). It has been modeled with one stated supertype and two attribute value pairs in a relationship group.
Figure 1: Stated view

The inferred view shows the logical definition of the concept. By using the stated relationships (for this concept and other concepts currently in the terminology), the classifier infers three defined proximal supertypes:

- Radiography of humerus (procedure)
- Computed tomography of upper arm (procedure)
- Computed tomography of bone (procedure)
Multiple potential primitive supertype concepts

Where more than one potential primitive supertype is identified for a concept, authors should check the primitive supertypes for subsumption of one or more other primitive supertypes. Any subsuming concept is not a PP supertype.

For example,

- There is more than one potential primitive supertype for 421095001 | Allergic disorder by body site affected (disorder)|. However, 64572001 | Disease (disorder)| is subsumed by 404684003 | Clinical finding (finding)|, therefore 64572001 | Disease (disorder)| is the proximal primitive supertype concept.

GCI-Modeled primitive supertypes

For information on the effect of GCIs on modeling primitive supertypes, see General Concept Inclusions (GCIs), GCI-Modeled Primitive Ancestor.
Intermediate Primitive Concept Modeling

Concepts that cannot be sufficiently defined by necessary conditions are called primitive concepts.

Primitive concepts cannot have subtypes automatically assigned by the classifier, unless a sufficient condition for that concept exists. Relevant concepts that are subtypes of a primitive concept in the taxonomy must be manually assigned an IS A relationship to that concept.

When a primitive concept is a child of one or more concepts and a parent of one or more concepts, it is known as an intermediate primitive.

For example,

- 116223007 |Complication (disorder)|

Without a stated IS-A relationship to the proximal primitive concept, |Complication (disorder)|, a concept will not classify as a subtype of |Complication|. Hence, all relevant subtypes will not be classified as complications.

Identifying all subtypes is important when creating a subset or when identifying relevant content during data retrieval. Therefore, when adding new concepts, potential primitive parents need to be identified and the IS_A relationship stated.

Consistent assignment of subtypes to intermediate primitive concepts is challenging. To find a possible intermediate primitive parent, it may be necessary to view the authoring form of several concepts that should be siblings of the new concept. Authors should also check for a possible intermediate primitive supertype among the descendants of the most proximate defined parent(s) under which the new concept would be expected to classify as an inferred subtype.

Given the manual burden that intermediate primitives impose, the creation of new intermediate primitive concepts in the international edition is prohibited unless:

- There is no other option and the concept is clinically necessary.
- The impact of adding the concept has been fully explored and understood.
- The impact is manageable and there is a management plan, including an extensional definition for the direct sub-concepts.

For the International Release, such requests are assessed case-by-case.

Relationship Group

A relationship group combines an attribute-value pair with none, one, or multiple attribute-value pairs in order to refine the meaning of a concept.

For example,
An attribute must be accompanied with a value in order to be used to model a concept. This page describes the grouping of attributes.

- A single relationship group containing only one attribute can exist.
  - When an attribute is restricted to a single group with no other attributes are allowed, the attribute is described as being "self-grouped".
- Multiple attributes may be grouped together in relationship groups, and multiple relationship groups may be created to sufficiently define concepts.
- When creating new concepts or revising existing ones, each attribute type included in a relationship group may only be present once, e.g. two Associated morphology attributes cannot be in the same relationship group.
- Relationship groups originated to add clarity to:
  - Clinical finding concepts which require multiple Associated morphology attributes and multiple Finding site attributes; and
  - Procedure concepts which require multiple Method attributes and multiple Procedure site attributes.

Figure 1: Stated view of 18876004 |Pain in finger (finding)| with the Finding site (attribute) and its value of Finger structure (body structure)
• Relationship groups are not limited to Clinical finding and Procedure concepts.
• There is no limit to the number of relationship groups that may be added to a concept.

**Modeling**

As with all authoring activities, grouping of attributes is performed in the stated view.

**Ungrouped attributes**

An attribute that is not in a relationship group is considered to be in a group on its own. When attributes are not grouped, their meanings are interpreted separately. For example, in the following diagram, the Associated morphology is Hemorrhage, and the Finding site is Uterine structure. However, it cannot be interpreted that the site of the Hemorrhage is the Uterine structure because the two attributes are not grouped.

![Figure 2: Inferred view of self-grouped attributes values of Hemorrhage (morphologic abnormality) and Uterine structure (body structure)](image-url)
When the attributes are grouped, the relationships imply meaning towards each other. To continue the example above for 44991000119100 | Abnormal uterine bleeding (disorder), the following diagram shows the Associated morphology of Hemorrhage and the Finding site of Uterine structure in a relationship group together. The grouping can be interpreted that the finding site of the hemorrhage is the uterine structure.

Figure 3: Inferred view of grouped attribute values of Hemorrhage (morphologic abnormality) and Uterine structure (body structure)

Note the difference in the inferred parents between the self-grouped versus grouped attributes. This is explained in more detail below.

Impact of relationship grouping on inheritance

Relationship groups refine inheritance, i.e. a grouped set of attributes is more specific than the same attributes that are not grouped. This is important when considering subsumption. The following diagrams demonstrate the impact of grouping or failing to group consistently using the concepts 50434004 | Excision of lesion of aorta (procedure) and one of its supertypes, 63296004 | Excision of aorta (procedure).
The meaning of the supertype concept, 63296004 | Excision of aorta (procedure) | (where the relationships are grouped) is interpreted as a procedure with an excision on the aortic structure. This is because 405813007 | Procedure site - Direct (attribute) and 260686004 | Method (attribute) are grouped.

Figure 4: Inferred view of Excision of aorta (procedure) with grouping of attributes

In the following diagram, the more general supertype concepts, 65801008 | Excision (procedure) | and 118809006 | Procedure on aorta (procedure) | are the proximal supertype concepts.

50434004 | Excision of lesion of aorta (procedure) | is a logical subtype of 63296004 | Excision of aorta (procedure) |. However, the attributes of the concept 50434004 | Excision of lesion of aorta (procedure) | are not grouped. Thus, the classifier interprets the definitions as non-related and 50434004 | Excision of lesion of aorta (procedure) | is not inferred as a subtype of 63296004 | Excision of aorta (procedure) |. This is because the attributes in the subtype concept are not grouped, i.e. are not explicitly stated. From a machine-processing perspective, each attribute is
considered a group on its own; i.e. there is an excision, but nothing else is known about the excision. This results in the concept, 63296004 | Excision of aorta (procedure), being interpreted more broadly.

Figure 5: Inferred view of Excision of lesion of aorta (procedure) without grouping of attributes

In the following diagram the attributes of the concept 50434004 | Excision of lesion of aorta (procedure) are grouped. An author that explicitly states that the excision is of a lesion found in the aortic structure, by grouping the attribute-value pairs, provides the necessary information for the classifier. This enables 50434004 | Excision of lesion of aorta (procedure) to be inferred as a subtype of 63296004 | Excision of aorta (procedure).
Figure 6: Inferred view of Excision of lesion of aorta (procedure) with grouping of attributes

Same attributes in separate relationship groups

Each relationship group should only contain one instance of an attribute. This is because two of the same attributes in a relationship group is not the same as one attribute with one target value that captures the combined meaning of the target values, as illustrated in the following diagram.

Two Finding site attributes are required to support the location of 53627009 [Closed fracture of radius AND ulna (disorder)]. Each 363698007 [Finding site (attribute)] and its respective target value are placed in a relationship group with the attribute 116676008 [Associated morphology (attribute)] with its target value of 20946005 [Fracture, closed (morphologic abnormality)].
Figure 7: Inferred view of Associated morphology (attribute) with its value of Fracture, closed (morphologic abnormality) in two separate relationship groups

Procedure hierarchy

In the 71388002 Procedure (procedure) hierarchy, a relationship group is usually a way of combining attributes about a particular method.

In the concept 302619004 Cholecystectomy and exploration of bile duct (procedure) within the following diagram, the relationship groups clarify that there is exploration of the bile duct and excision of the gallbladder. Without the
relationship groups, the appropriate relationships between the attributes would be unclear; i.e. the exploration of the bile duct versus gallbladder and the excision of the bile duct versus the gallbladder.

Figure 8: Inferred view of a Procedure hierarchy relationship group: combining attributes around Method (attribute)
Clinical Finding/Disorder hierarchy

In the Clinical finding/Disorder hierarchy:

- The Finding site (attribute) and Associated morphology (attribute) are always grouped when both are present and related.
  - When there is more than one Finding site (attribute) or Associated morphology (attribute), then more than one relationship group is required.
  - When the attributes Occurrence and/or Causative agent are stated and related to the Finding site and Associated morphology attributes, include them within that relationship group.
    - As in the following diagram, when the Causative agent (attribute) is an organism, the Pathological process (attribute) is also included in that relationship group, with the target value of either 441862004 | Infectious process (qualifier value) or 442614005 | Parasitic process (qualifier value).
  - If a concept has values for a Causative agent (attribute) and Finding site (attribute), but does not have a value for an Associated morphology (attribute) or Pathological process (attribute), combine the Causative agent (attribute) and Finding site (attribute) as usual. Concepts that only have Causative agent (attribute) and Finding site (attribute) in a role group are higher in the hierarchy and subsume those concepts that have a role group of Causative agent (attribute), Finding site (attribute), Associated morphology (attribute) and Pathological process (attribute).
  - The Interprets and Has interpretation attributes are always grouped together where both are present and related to each other. These two attributes and their values are often used in defining a Clinical finding concept by delineating the observation results or describing the analysis used to determine the observation. Interprets and Has interpretation attributes are not grouped with any other attributes.
  - The Finding method and Finding informer attributes are also grouped together where both are present and related to each other.

Relationship group clarification

A relationship group that uses the attributes Before, During, After, Due to, Clinical course, or Temporally related to are not grouped with another relationship group; these attributes are "self-grouped". This means when modeling concepts, authors place these attributes in a relationship group individually with no other attributes.

- Note: 726633004 | Temporally related to (attribute) only applies to perioperative procedures and to a limited number of clinical findings.
Figure 9: Stated view of a disorder hierarchy concept with Causative agent and Pathological process attributes in the same relationship group
Situation with Explicit Context hierarchy

For **413350009** |Finding with explicit context (situation)| concepts, the following four attributes are grouped:

- 408729009 |Finding context (attribute)|
- 246090004 |Associated finding (attribute)|
- 408731000 |Temporal context (attribute)|
- 408732007 |Subject relationship context (attribute)|

For example, 704008007 |No family history of asthma (situation)| IS A 243796009 |Situation with explicit context (situation)|,

- 408729009 |Finding context (attribute)|, 410516002 |Known absent (qualifier value)|
- 246090004 |Associated finding (attribute)|, 195967001 |Asthma (disorder)|
- 408731000 |Temporal context (attribute)|, 410511007 |Current or past (actual) (qualifier value)|
- 408732007 |Subject relationship context (attribute)|, 444148008 |Person in family of subject (person)|

For **129125009** |Procedure with explicit context (situation)| concepts the following four attributes are grouped:

- 408730004 |Procedure context (attribute)|
- 363589002 |Associated procedure (attribute)|
- 408731000 |Temporal context (attribute)|
- 408732007 |Subject relationship context (attribute)|

For example, 704503005 |Advice given about pelvic floor exercise (situation)| IS A 129125009 |Procedure with explicit context (situation)|,

- 408730004 |Procedure context (attribute)|, 385658003 |Done (qualifier value)|
- 363589002 |Associated procedure (attribute)|, 420227002 |Recommendation to (procedure)|
Observable Entity hierarchy

When defining |Observable entity (observable entity)| concepts, attributes are self-grouped. Each observable entity represents only one property being observed. For example, |Standing diastolic blood pressure (observable entity)| is represented using multiple attributes with each in its own relationship group.

Figure 10: Stated view of a concept from the Observable entity hierarchy with self-grouped attributes
Templates

In addition to the guidance found here in the Editorial Guide, please see information on the use of templates at SNOMED CT Modeling Templates and Description Patterns.

Templates are created by authors in an attempt to standardize the modeling, naming, case significance, etc. of certain subhierarchies of the terminology, though it is recognized that not every concept may conform to a prescribed pattern. The modeling approach may be difficult to apply in all cases, but domain-specific templates are being developed to ensure modeling consistency and accuracy.

When to create a template

No template is necessary if less than 50 concepts are affected. In cases of small numbers, check if existing templates can be generalized and/or look to add elements as optional rather than mandatory.

Domain Specific Modeling

SNOMED CT is arranged as a polyhierarchy. A hierarchy is defined as an ordered organization of concept codes linked together through IS A relationships. Concept codes are linked to their more general parent concept codes directly above them in a hierarchy. Concepts with more general meanings are usually located at the top of the hierarchy and then at each level down the hierarchy the meanings become increasingly more specialized.

Selected SNOMED CT attributes have a hierarchical relationship to one another known as attribute hierarchies. In an attribute hierarchy, one general attribute is the parent of one or more specific subtypes of that attribute. Concepts defined using the more general attribute can inherit concepts modeled with the more specialized subtypes of that attribute.

Domains

The following are the 19 domains arranged in alphabetical order.

The following subhierarchies do not have concept models:

- Environment or geographical location (environment / location)
- Organism (organism)
- Physical force (physical force)
- Qualifier value (qualifier value)
- Record artifact (record artifact)
- SNOMED CT Model Component (metadata)
- Social context (social concept)
- Special concept (special concept)
- Staging and scales (staging scale)

HRCM Attribute Tables

The pages that follow contain tables that are generated by the Human Readable Concept Model (HRCM). The tables contain Attribute Summaries for those domains with attributes, information on Group(ed), Cardinality, and In-group cardinality, and Range constraints. The HRCM tables in this guide only reflect the ranges for pre-coordinated concepts; there may be post-coordination values that are not reflected in the tables. All MRCM values for concepts can be viewed via the public MRCM browser at https://browser.ihtsdotools.org/mrcm.

SNOMED International creates precoordinated content in accordance with the MRCM. For postcoordinated content, extensions should review the MRCM. If the MRCM does not specify that a particular value is allowed for a given content type (e.g. using an observable entity value for |Component| in a postcoordinated expression), then it must not be used in that content type (e.g. postcoordinated expressions). The MRCM rules for postcoordination must be strictly followed. This is important for interoperability, being able to query the resulting content consistently, etc. However, the MRCM does provide the option for extensions to extend or adapt the rules in a
controlled way if required (see the last section of 6. Considerations). This includes expanding the ranges and/or adding new attributes where required. This needs to be done carefully to ensure consistency and data integrity between editions.

There are special cases in the MRCM where an attribute may have two rows. This situation is caused by a new cardinality rule: a row for existing/legacy SNOMED CT content and a row for newly created content. The row that is applicable to new content will be marked by a "[New]" notation. See MRCM Maintenance Process.

### Modeling: precoordination patterns

SNOMED CT relies on the rules for usefulness to avoid excessive precoordination (see Scope section of Editorial Guide).

Approved precoordination patterns have been created and are available at: Pre-coordination Naming Patterns Project. For additional information about the fields used in precoordination, see: What the fields in the Pre-coordination Naming Patterns JIRA Project mean.

### Body Structure

<table>
<thead>
<tr>
<th>Definition</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Includes Anatomical structures and Morphologic abnormalities (subtype of body structure)</td>
<td>Body structure</td>
</tr>
<tr>
<td></td>
<td>• 38033009 Amputation stump (body structure)</td>
</tr>
<tr>
<td></td>
<td>• 91134007 Mitral valve structure (body structure)</td>
</tr>
<tr>
<td></td>
<td>Morphologic abnormality</td>
</tr>
<tr>
<td></td>
<td>• 189955008 Biopsy wound (morphologic abnormality)</td>
</tr>
<tr>
<td></td>
<td>• 31470003 Adenosarcoma (morphologic abnormality)</td>
</tr>
<tr>
<td></td>
<td>Cell</td>
</tr>
<tr>
<td></td>
<td>• 25029008 Agranular white blood cell (cell)</td>
</tr>
<tr>
<td></td>
<td>• 57184004 T lymphocyte (cell)</td>
</tr>
<tr>
<td></td>
<td>Cell structure</td>
</tr>
<tr>
<td></td>
<td>• 4897009 Cell membrane, prokaryotic (cell structure)</td>
</tr>
<tr>
<td></td>
<td>• 36229000 Entire axon (cell structure)</td>
</tr>
</tbody>
</table>

The body structure domain includes anatomical structures, as well as morphologic abnormalities, as follows:

- Body structure (body structure)
  - Anatomical or acquired body structure (body structure)
  - Anatomical organizational pattern (body structure)
  - Anatomical site notations for tumor staging (body structure)
  - Body structure, altered from its original anatomical structure (morphologic abnormality)
  - Nonspecific site (body structure)
  - Normal anatomy (body structure)
  - Topography not assigned (body structure)
  - Topography unknown (body structure)

### Tumor staging

Concepts under 258331007 |Anatomical site notations for tumor staging (body structure)| require review and reallocation.
Body Structure Attributes Summary

When authoring in this domain, these are the approved attributes and allowable ranges. They are from the Human Readable Concept Model (HRCM).

<table>
<thead>
<tr>
<th>Domain Information for 123037004</th>
<th>Body structure (body structure)</th>
</tr>
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<tbody>
<tr>
<td>Domain Constraint</td>
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<td>Parent Domain</td>
<td>-</td>
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<tr>
<td>Proximal Primitive Constraint</td>
<td>&lt;&lt; 123037004 Body structure (body structure)</td>
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<tr>
<td>Proximal Primitive Refinement</td>
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</table>

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<th>Body structure (body structure)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Attribute</td>
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<tr>
<td>733928003 All or part of (attribute)</td>
<td>0</td>
</tr>
<tr>
<td>733931002 Constitutional part of (attribute)</td>
<td>0</td>
</tr>
<tr>
<td>733933004 Lateral half of (attribute)</td>
<td>0</td>
</tr>
<tr>
<td>774081006 Proper part of (attribute)</td>
<td>0</td>
</tr>
<tr>
<td>733930001 Regional part of (attribute)</td>
<td>0</td>
</tr>
<tr>
<td>733932009 Systemic part of (attribute)</td>
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</tr>
</tbody>
</table>

<table>
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<tr>
<th>Domain Information for 91723000</th>
<th>Anatomical structure (body structure)</th>
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</thead>
<tbody>
<tr>
<td>Domain Constraint</td>
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<tr>
<td>Parent Domain</td>
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</thead>
<tbody>
<tr>
<td>Attribute</td>
<td>Grou ped</td>
</tr>
<tr>
<td>272741003 Laterality (attribute)</td>
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</tr>
</tbody>
</table>
Anatomical Concept Model

The Structure-Entire-Part (SEP) model

SNOMED CT uses a structure-entire-part triple, known as the SEP triple, to represent anatomical structures. The following Relationships provided a way for the anatomy in CTV3 to be mapped to RT:

The SNOMED CT anatomy hierarchy differentiates classes of entire anatomical entities from classes of parts of entire anatomical entities.

**Entire concept**: Denotes a class that is instantiated by entire anatomical entities of some kind: entire heart is instantiated by all individual hearts.

**Entity Part concept**: Denotes a class that is instantiated by all anatomical entities that are a proper part of some entity of a given kind: heart part is instantiated by all entities that are a proper part of some heart, e.g. my mitral valve, your right ventricle, Joe’s sinus node. Heart part is not instantiated by any heart.

**Entity Structure concept**: Subsumes both the related Entire and Part concepts. Consequently, it denotes a class which is instantiated by anything that instantiates either the Entire or the Part. For instance, Heart structure is instantiated by my heart, my mitral valve, your heart, your right ventricle, Joe’s sinus node, Joe’s heart, etc.

The code named Liver structure in CTV3 is equivalent to Liver structure in the diagram above. Both the CTV3 code for Liver structure and the SNOMED RT code for Liver are interpreted to mean Some or all of the liver. Site attributes (PROCEDURE SITE, FINDING SITE) will usually take the value liver structure rather than entire liver, since typically the site of a liver disorder or procedure on the liver is not necessarily the entire liver.

Purpose of the Structure concept
Adding the Entity Structure codes is a convenience to assist with the logic-based aggregation of references to the entity or its parts. The implication of this view is that the E of the SEP triple is the code that should be regarded as the one that represents the real anatomical entity that is named.

For example, the code for entire liver is the one that should correspond to the code for liver in the Foundational Model of Anatomy (FMA). The subtype hierarchy for entire liver fits much better with the FMA hierarchies, and indeed it might be possible to completely reconcile SNOMED’s non-Structure components with FMA anatomy.

A database has been developed that categorizes codes in the physical anatomical entity hierarchy according to their status as S structure, P Part or E Entire, and provides the corresponding S and P code for each E code. This should provide some value to implementers. It can help with navigation, coordination with formal ontologies of anatomy, and selection of codes for postcoordination.

Conventions for merging concepts from SNOMED RT and Clinical Terms v.3

Where there were two concepts with the same name, the SNOMED RT code was to become the S code, and the CTV3 code was to become the E code. There are still instances of unrecognized pairing of the RT-CTV3 S-E pair, where neither codes FSN has been changed according to the naming conventions in this document. When these unmatched pairs are identified, it is our practice to change the FSNs accordingly, and to make the E code have a subtype IS-A link to the S code.

S concepts without a corresponding E concept

Some S codes do not currently have a corresponding E code subtype, and there was no policy that required that such E codes be created during the merger of SNOMED RT and CTV3. However, it is likely that such a policy will be enforced in the future.

S Structure codes can subsume entities other than E or P

The SEP triple may give the impression that all S codes have exactly two children, one E and one P, with all of the remaining descendants placed under P. Again, in the past this degree of modeling consistency was not always followed. Some codes were purposely made subtypes of the S that are not strictly part of the corresponding E.

For example, perirenal tissue is a kidney structure but not a part of the kidney. It is used to define perirenal abscess so that it is subsumed by renal abscess. While a perirenal abscess is not strictly within the substance of the kidney, it is still considered a kind of renal abscess, and the S anatomy hierarchy is used to support this inference.

This policy has introduced undesirable variation and arbitrariness into the terminology, and future revisions will seek to eliminate these variations. Where a code is needed for a site that is really meant to extend to entities that are not part of any kidney, this will be made clear in the name, e.g. Structure of kidney and perirenal tissue.

Countable vs non-countable E entities

The E code needs to be interpreted with care when the name refers to entities that do not have the property of identity, meaning that they are not countable wholes, or could be interpreted as non-countable. In this circumstance, the interpretation of E means some portion of the thing being named.

For example, tissue and types of tissue such as fascia, muscle, tendon, bone tissue, connective tissue, skin, mucosa/mucous membrane, nerve tissue, etc. Muscle, tendon, bone and skin can identify a type of tissue as well as an individual organ of that type. Bone tissue has no identity, but a particular bone does have identity.

To use skin as the archetypal example, the E code for skin of finger means a portion of the skin of a finger, so all of its subtypes must also be portions of skin. The S code for skin of finger then has a subtype P which would mean proper part of a portion of skin of finger. This admits subtypes that are not kinds of skin, but may be parts of skin, including layers, e.g. epidermis of finger (meaning a portion of epidermis of finger) could be a proper part of a portion of skin of finger.

Tissues, layers, membranes: portions
We regard the $E$ code for $x$ tissue, $x$ layer to have the meaning *portion of $X$ tissue*, and therefore regional subdivisions of tissue types are direct subtypes.

For example, transitional epithelium of urinary tract, as an $E$ kind of code, should be a supertype of transitional epithelium of urinary bladder. The reason is that (portion of) transitional epithelium of urinary bladder is a kind of (portion of) transitional epithelium of urinary tract.

We also deal with layers the same way.

For example, we regard serosal layer and serosa tissue as meaning the same thing, since all serosal tissue is configured as a layer, and it can’t be a serosa without being a layer; and their $E$ codes mean portion of serosal layer or portion of serosal tissue.

As another example, layer of retina would be a supertype of nerve fiber layer of retina, and also a supertype of retinal epithelium, where retinal epithelium represents a portion of the epithelium of the retina and is therefore a kind of (portion of) a layer.

Groups

The identity/countability issue extends to a problem differentiating groups of entities from one of the group.

For example, consider $x$ = *lymph node group*, $y$ = *lymph node*. In this case, the group should be linked to the member via an appropriate *Relationship* (not yet in SNOMED CT), such as has-member. In those cases where $y$ is always necessarily a member of group $x$, it could be linked via a member-of *Relationship* (also not yet in SNOMED CT).

What does part of mean?

There are several possible ways of interpreting *part of*. In SNOMED CT, *A part of B* means that in normal anatomy, the entire structure $A$ is structurally included in $B$. Another way of saying it is that $A$ is part of $B$ if there is no part of $A$ that is not also part of $B$.

For example, the humerus is not part of the shoulder region, because the distal humerus is part of the humerus, and the distal humerus is not part of the shoulder region.

We do *not* use part of for non-anatomical meanings, such as grouping tests together in batteries, nor do we use it to indicate *Relationships* that are not strict anatomical inclusion.

Some recent work has begun to differentiate between part of that is reflexive (that is, an entity is in some sense a *part of itself*, much the same that a set can be viewed as a subset of itself), versus *proper part of*, where an entity cannot be a proper part of itself. For now, we regard part of *Relationships* as implying strict partonomy.

There is sometimes confusion about parthood as opposed to location.

For example, an embryo is not part of a mother’s body, but a kidney is. The anatomy section is composed mainly of canonical parts; but a few abnormal parts are included to permit them to be used as the location of tumors or injuries.

For example, a Meckels diverticulum is a body structure that is part of the small intestine, and it is also a morphological abnormality. Likewise some stomas and other post-surgical structures are considered part of the body. A transplanted liver or kidney would be considered part of the body, as a post-surgical structure, even though the transplanted organ is not genetically identical. Likewise transplanted bone marrow is part of the body.

Non-living implants and devices, and foreign bodies, on the other hand, are considered to be located in the body, but not part of the body.

✅ **Part of relationships**

For more information on part of relationships in the anatomy concept model, please see [Part of relationships (under development)](#).
Can the SNOMED CT relationships table be used to construct a part of hierarchy?

The currently distributed part of Relationships need to be much more extensively modeled and quality assured. At present they are not defining, that is, their Characteristic Type in the relationship file is additional, and, therefore, they do not affect the classifier behavior. A substantial amount of effort has gone into a draft of the updated part of Relationships; these will require review and approval before incorporation into the release. This will eventually result in the SEP triplet structures and part of relations being strictly paralleled. It is a matter of time to implement and quality assure the changes.

Why are part of relationships not defining?

The SEP structure, combined with the inference mechanism that is used with SNOMED CT, allows us to take advantage of anatomical Relationships to infer subsumption, IS_A Relationships between disorders, procedures, and other entities without reference to part of Relationships. The SEP structure also permits us to sufficiently define anatomical structures without reference to part of Relationships (making them necessarily true, but not among the necessary and sufficient conditions).

For example, the Structure of left hand can be sufficiently defined as a hand structure with laterality = left. This definition is sufficient. Converting the part of Relationships to have Characteristic Status = defining will require significant changes to the current model.

Entities with mass versus purely spatial massless entities

Points, lines, and surfaces can be considered to be massless. The FMA calls these immaterial. It is important to differentiate the codes/names for these entities from those that are intended to represent entities that have mass. At present, the concepts under anatomical spatial entity represent massless entities. Massless entities are not represented using the SEP model. It is conceivable that users may want to reference parts of a surface, and to enable this we would need to apply the SEP model to anatomical spatial entities, or else adopt defining part of Relationships.

Attributes used to define body structure concepts

Laterality
This attribute provides information on whether a body structure is left, right, or bilateral. It is applied only to bilaterally symmetrical body structures which exist on opposite sides of the body.

Unilateral
With the addition of lateralized content in the International Release, the need for unspecified unilateral concepts is removed, as well as potentially dangerous, if used directly in a patient record. Unilateral concepts will not be accepted and a review of existing content for potential inactivation is underway.

Laterality
Determine if an anatomy structure is lateralizable
The anatomy structures should only be pre-coordinated with laterality if they are lateralizable. All anatomy structures on the midline are not lateralizable. The Lateralizable Body Structure Reference Set has been developed and published as part of SNOMED International release. Please note that the refset requires an ongoing update for new additions.

In this guide, the lateralizable anatomy structures are divided into three types:

1. X
   - The type X represents any anatomy structure that is lateral to mid-sagittal plane.
2. Lateral half of Y
   - Y represents any anatomy structure that is symmetrical on the body’s middle-line which cannot be lateralized. However, the lateral half of Y can have laterality, e.g. right half of head.
3. {Part} of X or Y
   - {Part} represents any constitutional or regional part of anatomical structure of X or lateral half of Y.
Creation of lateralized anatomy structure
Both Left and Right structures must be added when adding the new pre-coordinated concepts for anatomy structure with laterality. Lateralization should not be routinely applied to Entire and Part of anatomy concepts unless the concept model requires such lateralized anatomy structure.

**Bilateral X anatomical structure (body structure)** must not be added. The concepts under 422525002 |Structure of bilateral paired structures (body structure)| are no longer in use in the international edition of SNOMED CT because bilateral concepts are defined by two role groups. However, these concepts may still be in use by extensions, in post-coordinations, or as values in information models. We would recommend users to review their usage and provide feedback to us. Users will be consulted before these concepts are ultimately inactivated.

**Term patterns for laterality**
Following are the most common term patterns for the representation of laterality for anatomy structures. Preferred terms should have the same description without a semantic tag. In descriptions in hierarchies such as clinical finding/disorder, procedure, observable entity, and situation with explicit context, 'structure of' can be omitted when body site is not a concept of Entire anatomy entity.

1. **Structure of + left/right + X or Structure of + left/right + half of Y**
   - Structure of left hand (body structure)
   - Structure of right lung (body structure)
   - Structure of left ring finger (body structure)
   - Structure of left renal artery (body structure)
   - Bone structure of left tibia (body structure)
   - Structure of right half of head (body structure)
   - Structure of right cerebral hemisphere (body structure)

2. **Structure of + {part} + left/right + X or Structure of + {part} + left/right + half of Y**
   - Bone structure of left hand (body structure) - bone structure is constitutional part of hand
   - Bone structure of phalanx of left ring finger (body structure)
   - Skin structure of right foot (body structure) - skin structure is constitutional part of foot
   - Skin structure of left index finger (body structure)
   - Bone structure of proximal right humerus (body structure) - proximal is regional part of humerus
   - Bone structure of shaft of left femur (body structure) - shaft is regional part of femur

3. **Special cases that are different from the above two common term patterns**
   - **a. Finger/toe is not sufficient** - Laterality should be placed before hand/foot when finger/toe or any part of a finger/toe is not further specified to individual digit, e.g. ring finger, index finger, great toe. For example,
     - Structure of finger of left hand (body structure) - correct
     - Structure of left finger (body structure) - incorrect

   - Bone structure of phalanx of finger of left hand (body structure) - correct
   - Bone structure of phalanx of left finger (body structure) - incorrect

   - Structure of nail of toe of right foot (body structure) - neither 'right nail' nor 'right toe' is correct
   - Note, hand/foot is redundant when an individually named finger or toe, e.g. index finger, great toe, has been specified in a description. For example,
     - Bone structure of phalanx of left index finger (body structure) - correct
     - Bone structure of phalanx of index finger of left hand (body structure) - it is not wrong, but 'hand' is redundant
     - Bone structure of phalanx of left index finger of hand (body structure) - it is not wrong, but 'hand' is redundant

   - **b. Structure of + {non lateralizable part} + left/right lung** Parts of symmetric structures should be symmetric. However, some body parts are exceptions. The laterality value in modeling is inherited from its lateralizable parent concept, e.g. left/right lung structure. Therefore, descriptions must clearly indicate such inheritances to avoid potential confusion. It is not accurate for a term such as
"right middle lobe" because there is no "left middle lobe". The following are examples for correct description pattern. Note: existing content has not been following the term pattern.

- Structure of middle lobe of right lung (body structure)
- Structure of apical segmental bronchus of right lung (body structure)

c. Structure of + {part} + left/right half of Y Leave the new concept as primitive if concept 'Structure of half of Y lateral to mid-sagittal plane (body structure)' does not exist. The new concept for 'Structure of half of Y lateral to midsagittal plane' should not be added until the policy is developed.

Skin structure of left half of face (body structure) - 'skin of left face' is not accurate because it missed the word half.

Concept modeling for laterality
The 'part of' relationship should not be used for concept modeling in anatomy. The laterality attribute should be the only attribute for the representation of laterality. The new concept model for anatomy has not been implemented, and the proximal primitive modeling style should not be applied.

For term pattern 1, the concept X or lateral half of Y should be used to fully define a concept with laterality.

For example,
For term pattern 2, the pre-coordinated concept (part) of X should be used to fully define the concept with laterality. Note: Concepts for which an identifier has not been assigned have been shown with an identifier of '1111111111'.

For example,

Please note that the concept modeling and utility of pre-coordinated concepts of 'structure of (part) of lateral half of Y' and 'structure of half of Y lateral to mid-sagittal plane' are still under investigation. Similar new anatomy concepts should not be added. The concept model should only use existing pre-coordinated concept (part) of lateral half of Y.

For example,
If pre-coordinated concept does not exist for '{part} of structure of half of Y' or 'structure of half of Y lateral to mid-sagittal plane', the concept can be defined by (part) of Y and the definition status should be primitive.

For example, concept |Structure of lateral half of lower back| does not exist, the concept 'Structure of left half of lower back' should be defined as a primitive concept. Two parent concepts are expected: 61379005 |Structure of left side of trunk (body structure)| and 37822005 |Lower back structure (body structure)|. The additional parent |Structure of left side of trunk (body structure)| is to ensure that any lateralized concept must be a subconcept of a lateralizable structure.

Anatomical Structure Naming Conventions

Naming Convention for SEP Model

1. FSN must include the word 'structure', 'entire' or 'part' for concepts that are following the SEP model.

For example,

- Liver structure (body structure)
- Entire liver (body structure)
- Liver part (body structure)

2. All descriptions for Entire concept must contain the word 'Entire'.

For example,
3. The word 'structure' can be omitted for synonyms.

For example,

<table>
<thead>
<tr>
<th>FSN</th>
<th>PT</th>
<th>SYN</th>
</tr>
</thead>
<tbody>
<tr>
<td>Entire liver (body structure)</td>
<td>Entire liver</td>
<td></td>
</tr>
<tr>
<td>Entire thumb (body structure)</td>
<td>Entire thumb</td>
<td>Entire first digit of hand</td>
</tr>
</tbody>
</table>

S concepts are usually named x structure (body structure) or structure of x (body structure). E concepts are usually named entire x (body structure) or x entire (body structure). P concepts are usually named x part (body structure) or part of x (body structure).

Plurals
Outside the anatomy section of SNOMED CT, plurals were primarily used as headers, while the individual concept names were singular. In the anatomy section, we have taken plurals to represent meaningful differences from their singular counterparts.

For example, Skin structure of all fingers in the FSN would mean more than one finger, while Skin of finger would not imply more than one.

Naming Convention for Digits of Hand and Foot

Fully specified names and preferred names should use proper names of digits of hand and foot. The names by order of digits can be added as optional synonyms. The order of fingers is different from the order of digits of hand because the thumb is not a finger in SNOMED CT. Therefore, the order of the finger should not be used to avoid potential confusion.

Naming convention for the structure of digits of hand

<table>
<thead>
<tr>
<th>FSN</th>
<th>PT</th>
<th>SYN</th>
</tr>
</thead>
<tbody>
<tr>
<td>Index finger structure (body structure)</td>
<td>Index finger structure</td>
<td>Second digit of hand</td>
</tr>
<tr>
<td>Little finger structure (body structure)</td>
<td>Little finger structure</td>
<td>Fifth digit of hand</td>
</tr>
<tr>
<td>Middle finger structure (body structure)</td>
<td>Middle finger structure</td>
<td>Third digit of hand</td>
</tr>
<tr>
<td>Ring finger structure (body structure)</td>
<td>Ring finger structure</td>
<td>Fourth digit of hand</td>
</tr>
<tr>
<td>Thumb structure (body structure)</td>
<td>Thumb structure</td>
<td>First digit of hand</td>
</tr>
</tbody>
</table>

Naming convention for the structure of digits of foot

<table>
<thead>
<tr>
<th>FSN</th>
<th>PT</th>
<th>SYN</th>
</tr>
</thead>
<tbody>
<tr>
<td>Great toe structure (body structure)</td>
<td>Great toe structure</td>
<td>First digit of foot</td>
</tr>
<tr>
<td>Second toe structure (body structure)</td>
<td>Second toe structure</td>
<td>Second digit of foot</td>
</tr>
<tr>
<td>Third toe structure (body structure)</td>
<td>Third toe structure</td>
<td>Third digit of foot</td>
</tr>
<tr>
<td>Fourth toe structure (body structure)</td>
<td>Fourth toe structure</td>
<td>Fourth digit of foot</td>
</tr>
<tr>
<td>Little toe structure (body structure)</td>
<td>Little toe structure</td>
<td>Fifth digit of foot</td>
</tr>
</tbody>
</table>

Anatomical Structure Modeling

Many terms that refer to body systems or tracts are used imprecisely in clinical practice and in medical publications. Ambiguities frequently arise with many of these terms. We have made the following definitions and distinctions in order to achieve internal consistency of the terminology. We recognize that it may not be possible to get universal
consensus for the names for each of these concepts. The goal is to be consistent and clear in defining the meaning of each concept and to allow users and system designers to present the terms that best reflect these meanings in their own implementations.

Body parts and regions

SNOMED CT uses the Foundation Model of Anatomy (FMA) definition of body part and body part subdivision for some concepts. For example, the joint regions discussed below are classified as body part subdivisions, since that is what is intended by the diseases and procedures that use these terms in their definitions. They are not body parts because they are defined, not by a set of bones, but rather by a particular joint and its surrounding structures. However, our interpretation of the word region is based on common usage and is intended as a three-dimensional structure, not the FMA two-dimensional definition of body region. In other words, these regions are not simply surface regions (skin), but also include the three dimensional underlying structures (subcutaneous tissues, bones, muscles, tendons, fascia, vessels and etc.).

Surface regions

Many concepts contain the phrase surface region. These could be interpreted as massless (immaterial) mathematical surfaces, but a clinical terminology would have no direct use for such meanings in clinical records. They could be interpreted as having mass (not immaterial), but the depth then is arbitrary. Should it be just skin deep, or should it include deeper layers of the surface? If only skin deep, the meaning of these concepts would overlap with concepts for skin regions. If deeper, the meaning would possibly be the same as the generic structure concepts.

Abdominal regions

The named regions of the abdomen are by tradition divided horizontally by the transpyloric plane and the interspinous plane, and vertically by the midclavicular plane. The lateral regions are therefore bounded above by a plane that is inferior to the ribs. In contrast, the flank is the lateral region of the abdomen bounded above by the ribs. Thus some parts of the hypochondriac regions, which are superior to the transpyloric plane but inferior to the ribs, would be considered also part of the flank. The hypogastric region is also sometimes called the pubic region.

Abdominal cavity, pelvic cavity

The term abdominal cavity has two meanings, one including the pelvic cavity, the other excluding it. Abdominal cavity structure includes both. Abdominal cavity proper excludes the pelvic cavity.

Organs, organ system subdivisions

The FMA definition of body organ is also used. Organs include individual bones, joints, muscles, arteries, veins, lymph vessels, nerves, and etc. Concepts that include groups of organs are frequently used in SNOMED CT. In most cases, these have been part of the subsumption hierarchy (IS A hierarchy) of the particular organ type, that is, they are kinds of organs.

For concepts that refer to the collection of organs (rather than organs in a collection), there is another concept that is a kind of, organ system subdivision. Many such collections do not yet have corresponding organ system subdivision concepts. The default is to interpret concepts as denoting organs, rather than organ system subdivisions.

<table>
<thead>
<tr>
<th>Collections of Organs with/without Organ System Subdivisions</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Organ</strong></td>
</tr>
<tr>
<td>Vertebra (bone of vertebral column)</td>
</tr>
<tr>
<td>Cervical vertebra</td>
</tr>
</tbody>
</table>

Inactivation

Most surface region concepts will be retired as ambiguous/possibly equivalent to their corresponding concepts that are clearly not immaterial, including x structure, entire x, and skin of X. Where the x structure codes do not currently exist, they will be created, without the surface region phrase.
Collections of Organs with/without Organ System Subdivisions

<table>
<thead>
<tr>
<th>Organ Description</th>
<th>Organ System Subdivision</th>
</tr>
</thead>
<tbody>
<tr>
<td>Third cervical vertebra</td>
<td>No corresponding</td>
</tr>
<tr>
<td>Bone of skull</td>
<td>Skull (subdivision of</td>
</tr>
<tr>
<td></td>
<td>skeletal system)</td>
</tr>
<tr>
<td>Bone of thoracic cage</td>
<td>Thoracic cage (subdivision of skeletal system)</td>
</tr>
<tr>
<td>Rib</td>
<td>No corresponding</td>
</tr>
<tr>
<td>Third rib</td>
<td>No corresponding</td>
</tr>
<tr>
<td>Right third rib</td>
<td>No corresponding</td>
</tr>
<tr>
<td>Quadriceps femoris muscle</td>
<td>No corresponding</td>
</tr>
<tr>
<td>Quadriceps femoris muscle, left</td>
<td>No corresponding</td>
</tr>
<tr>
<td>Vastus medialis muscle</td>
<td>No corresponding</td>
</tr>
</tbody>
</table>

Cell, tissue, organ

In general, organs are made up of tissue, and tissue is made up of cells. However, a cell is not necessarily part of tissue, and tissue is not necessarily part of a named organ.

Tree structured organs

Arteries, veins, nerves, and the bronchi form tree-like structures that distribute across multiple regions. Because of their size and links with other structures, they require slightly different modeling. FMA divides tree structured organs as: **organs with organ cavities** and **organs that are solid**.

Hollow tree organs

**Organ with organ cavity** has a subtype, **hollow tree organ**. The hollow tree organs are:

- Tracheobronchial
- Biliary
- Vascular
  - Arterial
    - Systemic arterial
    - Pulmonary arterial
  - Venous
    - Systemic venous (superior, inferior, and 4 cardiac trees)
    - Pulmonary venous (superior and inferior left and superior and inferior right)
    - Portal venous
  - Lymphatic (right lymphatic duct and thoracic duct)

Among the solid organs, there is one category, **neural**, that is tree-structured (see: Nervous system; neural tree).

⚠️ Laterality of digits

For information on laterality, see Anatomical Structure Naming Conventions section at Naming Convention for Digits of Hand and Foot and Laterality section at Laterality

⚠️ Unacceptable terms

_X disorder at Y level_ concepts from ICD-11, e.g. skin laceration of arm at wrist level (precedent are terms added from ICD-9) will not be added to the SNOMED International Release.
Cardiovascular System

Cardiac valves, normal and malformed
There are a number of concepts in the anatomy hierarchy that represent congenital cardiac malformations. This content was developed in cooperation with IPCCC (International Pediatric and Congenital Cardiac Code).

The following pairs of cardiac valve concepts do not represent the same thing and are siblings, not super- or subtypes, to each other:

- 11124005 | Atrioventricular valve (body structure) | vs. 279316009 | Atrioventricular (non-mitral, non-tricuspid) valve structure (body structure) |
- 91134007 | Mitral valve structure (body structure) | vs. 312523009 | Left (non-mitral) atrioventricular valve structure (body structure) |
- 46030003 | Tricuspid valve structure (body structure) | vs. 244344000 | Entire right (non-tricuspid) atrioventricular valve (body structure) |

Atrioventricular (non-mitral, non-tricuspid) valves represent body structures which were anatomically abnormal from the beginning of their development. They are not called mitral/tricuspid valve although they perform the same function as their normal counterpart would. They are also represented using the term *not morphologically mitral/tricuspid valve*.

For example,

- 459176007 | Abscess of right atrioventricular (not morphologically tricuspid) valve (disorder) | represents an abscess of the right atrioventricular valve that has been developed abnormally from the beginning vs. 431189009 | Abscess of tricuspid valve (disorder) |

For a normally developed mitral/tricuspid valve, the term *left/right atrioventricular valve* can be used interchangeably. They are true synonyms. However, they cannot be used for abnormally developed valves, i.e. left atrioventricular (non-mitral)/right atrioventricular (non-tricuspid) valves.

Systemic, pulmonary circulation
The *systemic circulatory system* is the combined arterial and venous circulation that begins where blood leaves the left ventricle and ends where blood enters the right atrium. It excludes the coronary circulation. The heart chambers are also considered part of the circulatory system.

The *pulmonary circulation* is the combined arterial and venous circulation that begins where blood leaves the right ventricle and ends where blood enters the left atrium.

Arterial
81040000 | Pulmonary artery structure (body structure) |: Any artery of the pulmonary circulation, i.e. arteries carrying unoxygenated blood from the heart to the lungs. They include the trunk, right and left branches of the pulmonary artery (which are within the mediastinum), and all of their branches (which tend to occur at or past the hilum and are therefore regionally within the lung). 
128260003 | Pulmonary artery within lung (body structure) |: Any artery of the pulmonary circulation that is regionally within the lung, the boundary being defined by the hilum.
45341000 | Structure of trunk of pulmonary artery (body structure) |: The main pulmonary artery (one of the great vessels that enter the heart) carrying blood from the right ventricle and dividing into right and left main pulmonary arteries (some dictionaries consider this synonymous with pulmonary artery).

Venous
430757002 | Structure of pulmonary vein great vessel (body structure) |: There are four pulmonary veins that enter the left atrium, two on each side. These are what is intended by the name *pulmonary vein (great vessels)* that enter the heart. In common usage, any vein that is part of the lung may be referred to as a pulmonary vein, but SNOMED CT has a separate concept: 122972007 | Pulmonary venous structure (body structure) |: This means any vein that drains the lung. A synonym is *vein of lung*. *Pulmonary veins* are veins of the lung, but *pulmonary vein and vein of lung* are not synonyms.
Central, peripheral, cerebrovascular systems

The term *central vascular* is not in common use. In fact, the term does not appear in SNOMED CT. However, the term *peripheral vascular* is very common, and therefore it requires a definition that (by default) sets the boundary between central and peripheral vascular systems.

The simplest definition of the peripheral vascular system is the vascular system that is not central; and then the central vascular system includes the pulmonary circulation, coronary circulation, cerebrovascular system, thoracic aorta, superior vena cava, inferior vena cava, and mediastinal blood vessels.

Peripheral vascular disease is often distinguished from cerebrovascular disease and coronary artery disease. These are the three major categories of diseases caused by problems in vascular circulation in general, and atherosclerosis, in particular. As a result of this clinical distinction, the cerebrovascular system is excluded from the peripheral vascular system.

*Cerebrovascular* is commonly defined in two ways: the blood vessels *in* the brain, or the blood vessels that *supply* the brain (including those within the brain). Because cerebrovascular disease includes extra-cranial occlusions of the vertebral and carotid arteries, we define the cerebrovascular system as those vessels involved in the supply and drainage of blood to the brain. Convention does, however, tend to exclude the innominate artery - which gives rise to the left common carotid and the arch of the aorta which gives rise to the right common carotid. Convention also excludes the subclavian arteries which give rise to the vertebral arteries.

**Common carotid artery, artery of neck**
The common carotid artery has a left and right component. The right common carotid artery has no thoracic portion (it arises from the brachiocephalic trunk behind the right sternoclavicular joint). The left common carotid artery has a thoracic portion (it arises from the arch of the aorta). Thus, the common carotid artery (not specifying laterality) is not exclusively an artery of the neck. This is because of the thoracic portion of the left common carotid artery. Then, artery of neck region includes the cervical part of left common carotid artery and all of the right common carotid artery.

**Intracranial, extracranial vascular system**
Some vascular trees are located wholly within the cranial cavity, but some (internal carotid; vertebral) cross the boundary between extra- and intra-cranial. Intracranial segments of such vascular trees must be individually identified as such, and the entire vascular tree must not be categorized as either extra- or intra-cranial.

### Tree-structured organs
See *Tree-structured organs* elsewhere, re: regional sections of venous and arterial tree organs.

### The word artery
The word artery has three different meanings. In modeling SNOMED CT concepts that refer to arteries, it is necessary to decide on a case-by-case basis which of these meanings is intended.

<table>
<thead>
<tr>
<th>Meanings</th>
<th>Notes</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>An arterial trunk: a single tube</td>
<td>The most common in clinical use.</td>
<td>A puncture wound of the femoral artery affects the femoral arterial trunk.</td>
</tr>
<tr>
<td></td>
<td>The meaning of the word <em>artery</em> in injuries and operations is clearly a single tube, the trunk of the named artery, or trunk of the named arterial branch.</td>
<td>A grafting into the popliteal artery is done into the popliteal arterial trunk.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Occlusions of arteries are located by naming the trunk where the occlusion occurs. Occlusions may affect circulation beyond the trunk, however, collateral circulation often mitigates the effects. Thus, it is incorrect to interpret <em>artery</em> to mean the entire subtree in any of these usages.</td>
</tr>
</tbody>
</table>
Meanings of artery

<table>
<thead>
<tr>
<th>An arterial tree organ</th>
<th>There are only two complete arterial tree organs (the systemic arterial tree arising at the aortic valve, and the pulmonary arterial tree arising at the pulmonary valve) that are readily named as such. They are seldom referred to by disorders or procedures.</th>
<th>NA</th>
</tr>
</thead>
<tbody>
<tr>
<td>An arterial trunk, plus all its branches</td>
<td>When modeling, it is challenging to differentiate when trunk vs. trunk plus branches is intended.</td>
<td>NA</td>
</tr>
</tbody>
</table>

⚠️ Artery

This clinical usage of *artery* varies from the definition of the FMA, which defines *artery* as a subdivision of an arterial tree (organ) which consists of branching sets of tubes (arterial trunks) that form a tree; together with other arterial trees (organ parts), it constitutes an arterial tree (organ). The FMA definition corresponds to the third meaning of *artery* above.

The word vein

The word vein has three different meanings. In modeling SNOMED CT concepts that refer to veins, it is necessary to decide on a case-by-case basis which of these meanings is intended.

Vein

<table>
<thead>
<tr>
<th>Meaning</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>A venous trunk</td>
<td>As with the clinical usage of the word <em>artery</em>, clinical usage of the word <em>vein</em> generally refers to the trunk and not the entire tree.</td>
</tr>
<tr>
<td>A venous tree organ</td>
<td>There are only eleven venous tree organs that are readily named as such.</td>
</tr>
<tr>
<td>A venous trunk, plus all its branches</td>
<td>When modeling, it is challenging to differentiate when trunk vs. trunk plus branches is intended.</td>
</tr>
</tbody>
</table>

⚠️ Vein

This clinical usage of *vein* varies from the definition of the FMA, which defines *vein* as a subdivision of a venous tree (organ) which consists of branching sets of tubes (venous trunks) that form a tree; together with other venous trees (organ parts), it constitutes a venous tree (organ). The FMA definition corresponds to the third meaning of *vein* above.

Trunk of vein, vein as a tree structure

Because *trunks of veins*, not *venous trees*, have been used to organize the vein hierarchy, there are implications for regional classes.

For example, the internal jugular vein is a vein of the neck, but its entire venous tree extends into the head. The internal jugular vein venous tree is not strictly part of the neck, even though the internal jugular vein venous trunk is strictly part of the neck.

Tributaries are also modeled as direct tributaries of the trunk. A tributary of a named vein is part of the *venous tree* of the named vein, but not part of the *venous trunk* of the named vein. Some veins that are part of the *venous tree*, and therefore might be regarded as indirect tributaries, are not modeled as direct tributaries of the *trunk of the vein*. Direct tributary is the intended meaning of tributary.
**Digestive System**

*Digestive tract* is the same as alimentary tract, and includes the entire passage for food through the body, including mouth, oral cavity (both vestibule of mouth and cavitas oris propria), oropharynx, esophagus, stomach, duodenum, jejunum, ileum, colon, rectum, and anal canal.

*Digestive system* includes the digestive tract, as well as the associated organs of digestion, including tongue, teeth, salivary glands, liver, exocrine pancreas, gallbladder, and biliary tract.

*Gastrointestinal tract* has two meanings in common usage. One that does and one that does not include the esophagus. The usage that includes the esophagus would more correctly be named esophago-gastrointestinal tract. Endoscopists frequently use this meaning, even though it is contrary to some dictionary definitions and does not follow strict lexical interpretation (which does not include the esophagus).

- **Upper gastrointestinal (GI) tract.** When describing upper GI bleeding and upper GI radiographic and endoscopic procedures the upper GI tract includes the esophagus, stomach, and duodenum. The upper GI tract does not include the more restricted *stomach-intestine* entity.

- **Lower gastrointestinal (GI) tract.** When describing lower GI bleeding, lower GI radiographic and endoscopic procedures, and lower GI output from ileostomies and colostomies, the lower GI tract includes the jejunum, ileum, cecum, colon, rectum and anal canal. The ligament of Treitz may be used as the division between upper and lower GI tracts (and the division between the duodenum and jejunum).

  Also, since the upper GI tract is said to end at the duodenum-jejunum junction, and there is no concept meaning middle GI tract, the jejunum can be inferred to be in the lower GI tract.

---

**Upper aerodigestive tract**

The SNOMED CT concepts 119253004 |Upper aerodigestive tract structure (body structure)| and 361922007 |Entire upper aerodigestive tract (body structure)| have the meaning based on the following reference: *Cancers of the upper aerodigestive tract constitute approximately 4% of all malignancies. These include cancer of the lip, tongue, major salivary glands, gums and adjacent oral cavity tissues, floor of the mouth, tonsils, oropharynx, nasopharynx, hypopharynx and other oral regions, nasal cavity, accessory sinuses, middle ear, and larynx (Upper aerodigestive tract cancers, Cancer 1995 Jan 1;75 (1 Suppl): 147-53). This definition matches the tumors included in the *CAP Cancer Checklist* for upper aerodigestive tumors. The esophagus, or at least the cervical esophagus, may be included, but not in SNOMED CT.

Biliary tract

Biliary tract includes the gallbladder, intrahepatic and extrahepatic bile ducts, and common bile duct. It does not include the liver. SNOMED CT uses *biliary system* as a synonym for *biliary tract*. SNOMED CT has another concept that does include the liver, 732049009 |Entire liver and biliary system (body structure)|.

Mouth

Mouth has several different meanings including mouth region, oral region of the face, and rima oris.

Mouth region includes structures surrounding the oral cavity, as well as structures of the oral region of the face.

**Modeling**

Use *mouth region* for most disorders with a finding site of *mouth.*
Oral region of face includes the skin and subcutaneous tissue of the lips and perioral region, the orbicularis oris muscle, and the vessels and nerves in these structures.

Rima oris is the opening of the mouth.

Tongue
The four regional parts of the tongue are the ventrum (inferior surface), dorsum, root, and body. The root of the tongue is the posterior third, the dorsal surface of which forms the anterior wall of the oropharynx. The root of the tongue rests on the floor of the mouth. The nerves and vessels that supply the intrinsic muscles of the tongue traverse the root of the tongue.

Ear
The ear includes the external, middle and inner ear. The external ear has two main parts, the auricle (also called the pinna) and the 84301002 External auditory canal structure (body structure). The external auditory canal has the synonym external auditory meatus. The external auditory meatus is not just the external opening of the canal, but rather the canal extending to the ear drum (42859004 | Tympanic membrane structure (body structure)). The 61671002 Structure of internal acoustic meatus of temporal bone (body structure) | SYN, internal auditory canal, is not part of the ear. As described in the FSN, it is an opening in the temporal bone, and is primarily a nerve conduit that anatomically parallel to the external auditory canal.

Endocrine System
The endocrine system is composed of the endocrine pancreas, pineal body, paraganglia, paraaortic bodies, parathyroid glands, endocrine ovaries, endocrine testes, adrenal glands, pituitary gland, thyroid gland, juxtaglomerular apparatus of the kidneys, and some diffuse neuroendocrine structures. Certain parts of the thymus produce endocrine hormones, but the thymus itself is not part of the endocrine system.

Eye
Choroid
Both subchoroidal and suprachoroidal refer to the same potential anatomic space between the choroid and the sclera. The term lamina subchoroidea of choroid is the same as the lamina suprachoroidea.

⚠️ Suprachoroidal hemorrhage
In the literature, the term massive suprachoroidal hemorrhage is replacing expulsive hemorrhage and subchoroidal hemorrhage.

Retinal vein
There is not a vein actually named retinal vein. However, SNOMED CT has concepts with the phrase.

For example,
- 85003000 | Structure of retinal vein (body structure) | has the synonym retinal vein.
- 280927000 | Entire central vein of the retina (body structure) | has the synonym entire central retinal vein.

Orbital region
371398005 | Eye region structure (body structure) | has a synonym of orbital region structure which subsumes bony orbit, entire eye, and ocular adnexa.

Genitourinary system
The genitourinary system includes the entire urinary system, as well as the genital system. The genital system includes internal genital organs and external genitalia.

Urinary system/tract
The urinary system includes the organs that form and excrete urine, the kidneys, ureters, bladder, and urethra. The male urinary system includes the prostatic urethra (since it is a male urinary outflow structure).
In common usage, *urinary system* and *urinary tract* are used interchangeably. However, in SNOMED CT, this is not the case, i.e. they are not synonyms. The two concepts are: 122489005 | Urinary system structure (body structure) and 431938005 | Structure of urinary tract proper (body structure).

**Urinary tract proper**
The urinary tract proper includes the organs involved in the excretion of urine including the renal pelvis (but not the rest of the kidney), ureters, bladder, and urethra. It is used for disorders affecting the flow of urine (as opposed to its formation) or the urothelium, the lining of the urinary tract.

For example,
- 41368006 | Disorder of urinary tract proper (disorder)
- 249273002 | Finding of urinary tract proper (finding)
- 7163005 | Urinary tract obstruction (disorder)
- 255150000 | Carcinoma in situ of urinary tract proper (disorder)

**Upper urinary tract**
The upper urinary tract consists of the kidneys and the ureters (to the juncture with the bladder). Since upper urinary tract infections include kidney infection, the upper urinary tract must include the kidney.

**Upper urinary tract proper**
The upper urinary tract proper is the part of the urinary tract proper. It includes only part of the kidney, the renal pelvis, and the ureters.

For example,
- 25990002 | Renal pelvis structure (body structure) has a parent, 431491007 | Structure of upper urinary tract proper (body structure).

**Lower urinary tract**
The lower urinary tract, 19787009 | Lower urinary tract structure (body structure), is the urinary system below the junction of the ureter with the bladder. It consists of the bladder and urethra. Lower urinary tract and lower urinary system are the same. The male and female specific components are located under male urinary outflow structure and female urinary outflow structure, respectively.

**Obstetric and gravid**
*Obstetric and gravid* body structures should not be added in SNOMED CT. *Obstetric* is a context for a disorder, procedure, or medical specialty that is applied to a body structure during pregnancy, childbirth, or the postpartum period. The context does not change the body structure.

**Prostate lobes**
The *posterior lobe* of the prostate is described in newborns but does not persist in the adult. 113295002 | Structure of lobe of prostate (body structure) includes three lobes, left and right lateral, and medial.

**Integumentary System**
**Skin, skin-associated mucosa**
This is an example of a body structure that is used to group related terms. The concept 707861009 | Structure of skin and/or skin-associated mucous membrane (body structure) intentionally employs *disjunction* (inclusive Or). It includes structures in the deep layers, but excludes non-skin mucosal epithelium, e.g. bronchial, gastrointestinal, and genitourinary sites of squamous cell neoplasms. The 400199006 | Structure of skin and/or surface epithelium (body structure) concept is used to represent the sites of these neoplasms.

Skin and/or skin-associated mucosa is intended for use in dermatology. It is not intended to subsume all mucosal structures, which are under Mucous membrane structure (body structure).
Diseases of the skin
For the meaning of diseases of the skin, refer to the draft of ICD-11: Diseases of the skin incorporate conditions affecting the epidermis, its appendages (hair, hair follicle, sebaceous glands, apocrine sweat gland apparatus, eccrine sweat gland apparatus and associated mucous membranes (conjunctival, oral and genital), the dermis, the cutaneous vasculature and the subcutaneous tissue (subcutis).

Skin regions, skin of <named body part>
Since the phrase skin of finger can mean some or all of the skin of finger (if interpreted as a structure, rather than entire in the The StructureEntirePart (SEP) model, we could use IS-A to represent the relationship between skin of finger and skin of hand. Thus, skin of finger IS-A skin of hand, IS-A Skin structure of upper extremity, IS-A skin region. The word region is not used in all of these names, because it may refer to the entire region or a part of a region.

Scalp
Formal definitions of scalp include layers beneath the skin. Therefore we make a distinction between 41695006 | Scalp structure (body structure) and 43067004 | Skin structure of scalp (body structure).

Soft tissue
There are at least three different use cases and meanings, and thus categories, for the phrase soft tissue. They include:

- **Tumors.** Soft tissue gives rise to similar types of neoplasms of mesenchymal stem cell origin, generally called soft tissue neoplasms. This accounts for the inclusions/exclusions of the category. Non-neoplastic masses arising in soft tissue are included in the WHO Classification of Soft Tissue Tumours.
  - For tumors, soft tissue is defined as non-epithelial extraskeletal tissue of the body, exclusive of the mononuclear phagocyte system, glia, and supporting tissue of various mesenchymal organs. Other explicit inclusions are: fibrous tissue, fascia, ligaments, tendons, tendon sheaths, synovia, bursae, skeletal muscle, smooth muscle, fatty tissue, adipose tissue, blood vessels, lymph vessels, peripheral nerves, sympathetic and parasympathetic nerves, and ganglia, as well as subcutaneous tissue. Skin, skeletal cartilage, pleura, and the pericardium, peritoneum, central nervous system, endocrine glands, and viscera are excluded.

- **Sites of non-bone disorders and injuries of the limbs, head, neck, and body wall.** Skeletal cartilage, as well as all non-bone structures of the limbs, and subcutaneous tissue and fat are included. Skin and lymph nodes are not included. For the head, neck and torso, mononuclear phagocyte system, central nervous system, endocrine glands, viscera, and supporting tissues are excluded.

- **Structures identified in images.** Soft tissue include everything except for mineralized bone tissue and teeth.

Lymphatic, Immune, Hematologic, Hematopoietic systems

Lymphatic system / 89890002 | Structure of lymphatic system (body structure)
Set of structures through which lymph flows. It includes 59441001 | Structure of lymph node (body structure) and 83555006 | Structure of lymphatic vessel (body structure). It supports the categorization of findings, disorders and procedures that relate to the flow of lymph.

Lymphoid system / 12249001 | Lymphoid system structure (body structure)
Set of structures with groups of lymphoid cells, including those in the intestines, marrow, liver, and other locations, and the lymph nodes, spleen, thymus, and tonsils and adenoids; excludes the lymph vessels. It supports categorization of lymphomas.

Immune system / 116003000 | Structure of immune system (body structure)
All of the lymphoid system, as well as the mononuclear phagocytic system; the immune system also includes cellular and sub-cellular components involved in cellular and humoral immunity.

Mononuclear phagocytic system / 127908000 | Mononuclear phagocyte system structure (body structure)
Collection of true macrophages, distributed widely in the body (splenic and lymphoid sinusoids, liver Kupffer cells, pulmonary alveolar macrophages, osteoclasts, macrophages in serous membranes, and microgliocytes); also endothelial cells that line hematopoietic tissues.

Dendritic cell system / 127909008 [Dendritic cell system structure (body structure)]
Collection of antigen-presenting cells, including the following: epidermal Langerhans, dendritic reticulum, and interdigitating. Class I histiocytoses (Langerhans cell histiocytosis) are disorders of the dendritic cell system.

Hematologic system / 414387006 [Structure of hematological system (body structure)]
Bone marrow, the lymphoid system, the hematopoietic system, and the terminal cells of all lineages of the hematopoietic system (red cells, white cells, platelets, histiocytes, plasma cells, etc). Disorders of the hematologic system do not necessarily include disorders of the hemostatic system, even though bleeding and thrombosis are usually categorized as hematologic.

Hematopoietic system / 57171008 [Hematopoietic system structure (body structure)]
Structures and cells responsible for erythropoiesis, granulocytopoiesis, monocytopoiesis, thrombocytopoiesis, and lymphopoiesis. Refers to the immature cellular elements that eventually form the cellular components of blood. The blood itself cannot be strictly part of the hematopoietic system, since this would cause all components of blood to be part of the hematopoietic system (including components like albumin, clearly not hematopoietic). SNOMED CT considers leukocytes, erythrocytes, and platelets the result of hematopoiesis, but not blood-forming, otherwise leukocytosis would become a disorder of hematopoiesis, whereas it can arise simply from a demargination of white cells following stress. SNOMED CT has a concept named 419333002 | Cellular component of blood (substance)]; note that platelets are not actually cells, but are cellular components.

Blood
The blood is not necessarily part of the cardiovascular system, nor is it necessarily part of the hematopoietic system. 87612001 | Blood (substance) | is a body fluid, not strictly part of either the hematopoietic or cardiovascular systems.

Regional lymph nodes of lungs
SNOMED CT has lymph node concepts per their anatomical locations, e.g. pulmonary, bronchopulmonary, tracheobronchial, tracheal, and esophageal) and concepts for node groups used for clinical staging of lung cancer, i.e. lymph nodes categorized into 14 stations.

Professional societies concerned with the clinical staging of lung cancer have developed at least three different nomenclatures for stations of lung-related lymph nodes. Even though the numbering of the stations is very similar, the inter-relationships between the various node groups are complex, particularly in stations 4 and 10, near the carina and hilar regions.

SNOMED International considers American Joint Committee on Cancer (AJCC) Station 10, hilar lymph node, bronchial lymph node, and bronchopulmonary lymph node as synonyms. The American Thoracic Society (ATS) Station 10R, the right tracheobronchial lymph node is not a subtype of tracheobronchial lymph node because its definition includes nodes covered by both lower paratracheal lymph node, (AJCC Station 4) and by the hilar lymph node (AJCC Station 10). SNOMED CT uses tracheobronchial lymph node as a supertype of both inferior tracheobronchial (subcarinal) and superior tracheobronchial (a subset of lower paratracheal).

Musculoskeletal System
Skeletal system, bony skeleton
The skeletal system (systema skeletal in Nomina Anatomica) includes bones and cartilage. The bony skeleton includes bones only. The vertebral column is part of the skeletal system, and includes the intervertebral discs (fibrocartilage). Individual vertebrae are part of the bony skeleton.
In ordinary usage, bone combines the meanings bone organ and bone tissue.

The 5 anatomical concepts related to bone are:

1. 3138006 | Bone (tissue) structure (body structure). Tissue type that makes up bones; a quantity of regular connective tissue consisting of osteocytes and related cells, the intercellular matrix of which is ossified; or any part thereof.
2. 90780006 | Entire bone (organ) (body structure). Individual bones, e.g. femur, tibia, ulna, scaphoid, lunate. An organ with cavitated parts; consists primarily of compact (cortical) and cancellous bone surrounding bone marrow cavities; also includes periosteum, endosteum (and, according to FMA, articular cartilage).
3. 118966000 | Skeletal system subdivision (body structure). Groups of bones, e.g. spine, skull, bony pelvis.
4. 128530007 | Entire bony skeleton (body structure). Pars ossea systematis skeletalis, bone part of the skeletal system.
5. 113192009 | Skeletal system structure (body structure). Entire skeletal system, including bones and cartilage.

Bone (tissue) is part of entire bone (organ); entire bone (organ) is part of skeletal system subdivision (system); skeletal system subdivision (system) is part of entire bony skeleton (body structure); and entire bony skeleton (body structure) is part of skeletal system structure (body structure). We can use Entire bone (system) to define aggregate concepts that involve bones.

Non-ossified bone
Bone organs are composed primarily of bone tissue, but there are some non-ossified parts. In particular, periosteum is clearly a part of a bone organ, but is not ossified tissue.

Bone marrow, marrow cavity
Bone marrow is contained within the marrow cavity, but it is not part of the bone organ. The (empty) marrow cavity is part of the bone organ. The bone marrow structure (body structure) is not a subtype of Bone structure (body structure).

Clinically, marrow disorders are not usually considered bone disorders, nor are marrow procedures considered bone procedures.

For example,

- Bone marrow disorders are not musculoskeletal disorders, but bone disorders are musculoskeletal disorders. Bone marrow transplants are not considered types of bone transplant.
- 60168000 | Osteomyelitis (disorder) is not the same as 44462005 | Osteitis (disorder).

Structure of (named bone), bone structure of (named bone)
To differentiate marrow, vessels, nerves, and periosteum from the actual hard tissue of bones, we differentiate structure of tibia from 12611008 | Bone structure of tibia (body structure). The bone marrow and other soft tissues of the tibia can then be categorized separately from the hard tissues. Bone marrow diseases are not considered musculoskeletal diseases, so bone marrow structures should not be placed in the bone (tissue) structure hierarchy.

Long bone, short bone
ICD does not use the standard anatomical definition of long bone.
For example, Benign neoplasms of long bones are distinguished from benign neoplasms of short bones; the bones of the hand are considered short bones. The anatomical definition of long bone cites the proportional relationship between length and width (length >> width). It is clear that metacarpals, metatarsals, and phalanges are included in the anatomical definition of long bone.

In order to accommodate the differences between anatomical definitions and classifications, SNOMED CT has anatomical groupings that correspond to the ICD groupings. Scapula, humerus, radius, or ulna and long bone of thigh or lower leg are used as the sites for grouper concepts that match ICD definitions and groupings.

Sternum, manubrium, body, xiphoid
The sternum is considered a bone organ. The manubrium, body, and xiphoid are parts of the sternum, classed as zones in the FMA.

Teeth, maxilla, mandible
Even though teeth are supported by the maxillary or mandibular bone, they are not part of the Bone structure of maxilla (body structure) or Bone structure of mandible (body structure). Teeth are part of the Bone structure of maxilla (body structure) and Bone structure of mandible (body structure).

Joints, joint regions
In many diseases and procedures, reference is made to areas of the body that may ambiguously imply either a joint or a region surrounding the joint. Some common ones are:

<table>
<thead>
<tr>
<th>Joint vs. Joint Region</th>
<th>Anatomical Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>70258002</td>
<td>Ankle joint structure (body structure)</td>
</tr>
<tr>
<td>74670003</td>
<td>Wrist joint structure (body structure)</td>
</tr>
<tr>
<td>85537004</td>
<td>Glenohumeral joint structure (body structure)</td>
</tr>
</tbody>
</table>

Shoulder girdle
272691005 | Bone structure of shoulder girdle (body structure). This concept is used to define diseases and procedures affecting bones in the shoulder region, i.e. proximal humerus, scapula, and clavicle. It is not a bone, but a bone structure, and is part of the shoulder region.

Intertarsal joint structure
27949001 | Intertarsal joint structure (body structure); SYN: Tarsal joint: This structure is part of a group of bones forming the tarsus or tarsal joint (ankle). The 27162001 | Talocalcaneonavicular joint structure (body structure) is the articulation between the talus (one of the seven bones of the ankle joint) and the other bones of the tarsus, and is what is meant by the rarely-used term talotarsal joint. The talocalcaneal joint is a synonym for the Subtalar joint structure (body structure). Dislocations of the subtalar joint usually involve the Subtalar joint structure (body structure). The subtalar and talonavicular joints constitute the talocalcaneonavicular joint.

Arm, leg, upper, lower, extremity, limb
The meaning of the words arm and leg may be misinterpreted.

- Arm may refer to the upper limb, but it may also refer to the upper part of the arm.
- Leg may refer to the lower limb, but it may also refer to the lower part of the leg.
- In common usage, leg is a synonym of lower extremity, and arm is a synonym of upper extremity.

In SNOMED CT,
- 53120007 | Upper limb structure (body structure) includes the shoulder, upper arm, forearm, wrist, and hand
- Upper arm is a synonym to 40983000 | Structure of upper extremity between shoulder and elbow (body structure).
- 61685007 |Lower limb structure (body structure)| includes includes the hip, thigh, lower leg, ankle and foot.
- Lower leg is a synonym to 30021000 |Structure of lower extremity from knee to ankle (body structure)|. Lower leg does not include the foot. *Stedman's Medical Terminology* defines lower leg as *the segment of the inferior limb between the knee and the ankle*.

The word *limb* appears in the FSN of the body structure, while the word *extremity* appears as a synonym. Therefore, when constructing an FSN for a new clinical finding concept, this precedent should be followed:

FSN: 61685007 |Lower limb structure (body structure)|
PT: Lower limb structure
Synonym: Lower extremity

Additional descriptions of *leg* and *arm* are permitted for concepts whose FSNs refer to *lower limb* and *upper limb* respectively.

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**External sources**

External sources, such as WHO Classifications, may have conventions for interpreting the meaning of phrases that contain the words *arm* and *leg*. These sources may be referenced to help determine the meanings of *International Classification of Diseases (ICD)* terms when mapping or completing other actions. ICD terms may differ from common usage and will not necessarily match SNOMED CT concepts.

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**Shoulder and hip regions, upper and lower limbs**

The shoulder region is part of the upper limb, and the hip region is part of the lower limb. This follows the general pattern used in the *Foundation Model of Anatomy (FMA)*. The *FMA* defines the upper limb as the free upper limb and the pectoral girdle (of which the shoulder region is part) and the lower limb as the free lower limb and pelvic girdle (of which the hip region is part). SNOMED CT has the concept 699617006 |Structure of free lower limb (body structure)|, i.e. the lower limb not including the pelvic girdle. There is not a concept for the *free upper limb*.

**Axilla**

The axilla is bound by the upper limb laterally and the thorax medially. It may be viewed as not strictly part of the upper limb or the thorax or it may be views as part of both. 91470000 |Axillary region structure (body structure)| is defined in SNOMED CT as being both an upper limb structure and a thoracic structure.

**Tendon**

A muscle may be considered an entire functional unit, including attachments to the skeletal system, or merely the contractile part of this unit. In clinical use, muscle is the contractile part only. The *FMA* definition implies that tendons should be considered part of their corresponding muscles, rather than organs in their own right. SNOMED CT models |Tendon structure| as a subtype of |Structure of muscle and/or tendon|. Muscle and tendon are two separate anatomical entities.

For example, the 61352006 |Structure of achilles tendon (body structure)| is not a 53451005 |Triceps surae muscle structure (body structure)| (gastrocnemius and/or soleus) muscle structure.

**Muscle function**

When modeling muscle categories according to their functions, assume they mean the function of the *entire muscle*, unless stated otherwise.

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**Unacceptable terms**

*X disorder at Y level* concepts from ICD-11, e.g. *skin laceration of arm at wrist level* (precedent are terms added from ICD-9) will not be added to the SNOMED International Release.

**Anatomy relating to the spine**

The terms 'spine' and 'vertebral column' in descriptions of conditions or procedures are often used loosely in clinical discourse but can relate to three different general anatomical concepts:

421060004 |Structure of vertebral column (body structure)| (synonym: Spinal column, ‘Backbone’, Spine)
- includes the bones of the spine and associated joints and ligaments
  289959001 [Musculoskeletal structure of spine (body structure)]
- includes the vertebral column plus muscles and tendons associated with the spine
  1141981001 [Structure of vertebral column region (body structure)]
- includes the musculoskeletal structure of spine plus spinal canal, meninges, spinal cord, roots and ganglia
  and immediate soft tissue including adjacent vascular structures where specified.

The last concept has been allocated the following definition: This volume includes the spinal column, its spaces and contents, notably the spinal canal, spinal meninges and spinal cord. It also includes the muscles associated with the spine within (and including) the anterior and posterior thoracolumbar fascia and its equivalent nuchal fascia in the neck. This region also includes the spinal ventral (anterior) and dorsal (posterior) roots, the dorsal root (spinal) ganglions and the spinal nerve trunks.

To emphasize the different volumes, cross-sections are shown below of (Figure 1) the vertebral column in yellow, and (Figure 2) the larger vertebral column region in amber color.

![Figure 1, Vertebral column](image-url)
From the preceding descriptions, it is clear that *Vertebral column region* is a broader concept for 'spine', and it should be used when a procedure or condition could involve not only the bony component of the vertebral column, but also the spinal cord, nerve root, muscle, bone, or joint of spine.

For example,

- MRI of thoracic spine (procedure) is modeled with 1141986006 |Structure of thoracic vertebral column region (body structure)|.
- Pain in sacrum (finding) is interpreted as being related to the more general notion of 1144746008 |Structure of sacral vertebral column region (body structure)|.

When a procedure or disorder is exclusively related to musculoskeletal structures of the spine, the 'musculoskeletal structure of spine' should be used.

For example,

- Cervical traction (procedure), Manipulation of the cervical spine (procedure), and Rotational deformity of cervical spine (finding) are all modeled with 297166009 |Structure of musculoskeletal system of cervical spine (body structure)|.

Where a procedure or condition only effects the bone, joint, or ligament component (and not the muscles or tendons directly) the 'vertebral column variant' is used for modeling.

For example,

- Benign neoplasm of lumbar vertebral column (disorder) and Kyphoplasty of fracture of lumbar spine using fluoroscopic guidance (procedure) are both modeled with 122496007 |Structure of lumbar vertebral column (body structure)|.
Junctional and combined segments of spine
The spine is traditionally divided into the following regions:

- Cervical
- Thoracic
- Lumbar
- Sacral
- Coccyx

The anatomy of these individual segments are structured in accordance with the description above.

The regions and joints between these identified segments are referred to as cervicothoracic, thoracolumbar, lumbosacral and sacrococcygeal. The meaning of these words are subject in common parlance and some literature to be ambiguous.

For example, ‘Thoracolumbar’ sometimes refers to the thoracic spine and lumbar spine, or alternatively, it is used to express the junction between the thoracic spine and lumbar spine.

To avoid false assumptions, the junction of spinal segments in SNOMED CT have been made explicit by including the word junction in descriptions. This avoids potential misinterpretation as to whether a word such as ‘thoracolumbar’ relates to both spinal segments or just the adjacent volume.

For example,

- The notion of 1145014005 |Structure of thoracolumbar junction of vertebral column (body structure)| is only used for modeling when it is explicitly stated in the target concept e.g. 281907005 |Fracture dislocation of thoracolumbar junction (disorder)|.

By contrast, in the circumstance where a dependent concept relates to the combination of two segments, e.g. 702487007 |CT of thoracolumbar spine (procedure)|, (FSN Computed tomography of thoracic and lumbar spine), the concept is modeled with two axioms, namely Structure of lumbar vertebral column region and Structure of thoracic vertebral column region.

The volume or extent of the junctional zones themselves are not defined consistently in the literature but most commonly relate to the junction between two segments and one vertebra above and below. So the convention used in the SNOMED CT anatomy hierarchy follows this guidance.

For example,

- 1145014005 |Structure of thoracolumbar junction of vertebral column (body structure)| includes the following concepts in its class:
  - 66794005 |Bone structure of L1 (body structure)|
  - 23215003 |Bone structure of T12 (body structure)|
  - 714833001 |Structure of intervertebral syndesmosis of T12 and L1 (body structure)|
  - 181879009 |T12/L1 facet joint (body structure)|

In addition, a further convention is required as to where segmental junctional joints belong, i.e. either to the cephalic or caudal segment. For instance, does the T12/L1 facet joint relate to the Thoracic spine joint structure or Lumbar spine joint structure?

The convention used (based on common criteria of spinal injuries) is that the junctional joints are included with the cephalic segment. So, in the case of the T12/L1 facet joint, it is included in the class of Thoracic spine joint structure (body structure).
Similarly, 8454000 |Lumbar spine joint structure (body structure)| subsumes:

- Intervertebral L5-S1 disc
- Structure of lumbosacral joint
- Structure of facet joint between L5 and S1

Nervous System

The nervous system has two parts, central and peripheral.

- The central nervous system, sometimes also called the *neuraxis*, consists of the brain and spinal cord. The pyramidal system is a subdivision of the central nervous system; the extrapyramidal system is part of the brain.
- The peripheral nervous system includes all neural structures outside the central nervous system.

The nervous system is also divided as: autonomic, somatic, and enteric.

- The autonomic system is further divided as sympathetic and parasympathetic. The autonomic system is not entirely a part of the peripheral nervous system, but the autonomic nerves are peripheral.

Nerve

The word *nerve* has multiple meanings according to the FMA:

- nerve trunk
- neural organ (trunk plus branches, excluding nuclei, ganglia, and roots)
- neural tree organ, including nuclei, ganglia, roots, etc.

A *neural tree organ* is defined in FMA as *a nonparenchymatous organ which has as its parts an aggregate of neurons (nuclei or ganglia) and their axons which are grouped into fasciculi by connective tissue to form elongated, cable-like structures that are arranged into a tree. A nerve, according to FMA, is defined as *a segment of a neural tree organ which has as its parts a nerve trunk and its branches; together with other nerves of the same tree, it constitutes a neural tree.* The neural tree structure includes:

- Cranial nerve
  - Complex cranial nerve-tract
- Spinal nerve
- Spinal accessory nerve (strictly neither cranial or spinal nerves)
- Peripheral nerve
- Autonomic nerve

*Nerve*, conventionally has two meanings:

- An anatomically distinct nerve trunk (without branches) that is identified in a dissection (e.g. the structure that student identifies when a pin is placed in the trunk of the vagus nerve, for instance located on the arch of the aorta)
- A larger anatomical entity which supports a related set of functions (e.g. all anatomical components of the vagus nerve that are necessary for it to execute its functions (e.g. when a student is asked which nerve is responsible for slowing the heart the answer, *the vagus nerve*, includes the vagal nucleus, as well as the trunk and branches of the vagus).
- Neural tree designates the second concept in order to distinguish it from the first which is only a part (subdivision of) the vagal neural tree.

A third meaning of nerve, defined by the FMA is: *Segment of neural tree organ which has as its parts a nerve trunk and its branches; together with other nerves of the same tree it constitutes a neural tree.*

For example,

- Chorda tympani, digastric branch of facial nerve, greater petrosal nerve, posterior cutaneous branch of posterior ramus of cervical nerve, superior lateral cutaneous nerve of arm.
- If one severs the facial nerve, the meaning refers to the trunk. But if one has facial nerve palsy, the meaning refers to the entire distribution of the nerve and the functions served by it.
Inactivation

There were several concepts with the phrase *x nerve and its branches*, interpreted as meaning *the entire nerve and its branches*. Therefore, *x nerve and its branches* would be a duplicate of *entire x nerve*, when we interpret *entire x nerve* as being a neural tree organ.

For example,

- *Entire facial nerve* is a neural tree organ, so there is no need for an additional concept called *facial nerve and its branches*.
- *Entire cranial nerve* is a neural tree organ and *structure of cranial nerve* is that organ or any part (or branch) thereof. Branches of the cranial and spinal nerves are segments of the neural tree organs from which they branch.

All concepts named *nerve x and its branches* were inactivated due to their ambiguity. There are MAY BE A links to *structure of nerve x*, and *entire nerve x*. Specifying *trunk of a nerve* requires a specific concept.

Supratentorial brain

Cerebrum may refer to the *supratentorial brain*, which is everything except the midbrain, medulla, pons, and cerebellum. In this interpretation, the telencephalon and diencephalon are in the cerebrum. On the other hand, cerebrum may only refer to the parts derived embryologically from the telencephalon, the cerebral hemispheres and the intercerebral commissure (corpus callosum and anterior commissure).

*Supratentorial brain* may be used for categorizing tumors and for designating the location of swelling that can result in herniation. The telencephalon and diencephalon (including thalamus, geniculate bodies, pineal body, habenulae, and hypothalamus) are definitely supratentorial. The upper part of the midbrain (mesencephalon) is also supratentorial. SNOMED CT excludes all midbrain structures from the supratentorial brain.

Respiratory System

Respiratory tract

321667001 [Respiratory tract structure (body structure)]. In SNOMED CT, *respiratory tract* has the same meaning as the Nomina Anatomica term *apparatus respiratorius*, which includes the structures through which air passes from the nares to the alveoli. The *oral cavity* is not included. In common usage, *respiratory system* may have the same meaning as *respiratory tract*; but not in SNOMED CT. *Respiratory system* does not mean the global respiratory system that might include the CNS components of breathing. *Pleura* are part of the lower respiratory system, but not a part of the lower respiratory tract.

Upper aerodigestive tract

This phrase has several meanings. The SNOMED CT concepts 119253004 [Upper aerodigestive tract structure (body structure)] and 361922007 [Entire upper aerodigestive tract (body structure)] have the meaning based on the following reference: *Cancers of the upper aerodigestive tract constitute approximately 4% of all malignancies. These include cancer of the lip, tongue, major salivary glands, gums and adjacent oral cavity tissues, floor of the mouth, tonsils, oropharynx, nasopharynx, hypopharynx and other oral regions, nasal cavity, accessory sinuses, middle ear, and larynx* (Upper aerodigestive tract cancers, Cancer 1995 Jan 1;75 (1 Suppl): 147-53). This definition matches the tumors included in the *CAP Cancer Checklist* for upper aerodigestive tumors. The esophagus, or at least the cervical esophagus, may be included, but not in SNOMED CT.

Upper respiratory tract

58675001 [Upper respiratory tract structure (body structure)] includes the nasal cavity, paranasal sinuses, nasopharynx, oropharynx, and larynx

Lower respiratory tract

82094008 [Lower respiratory tract structure (body structure)] includes the tracheobronchial tree (from the trachea through the terminal bronchioles) and the lungs, including the alveolar respiratory tract (which extends from the respiratory bronchioles to the alveoli).

Lower respiratory system

400141005 [Lower respiratory system structure (body structure)] includes the lower respiratory tract and the pleura.
Interarytenoid fold or larynx
The interarytenoid fold forms part of the inlet of the larynx. The fold has two surfaces, one forming part of the wall of the supraglottic larynx, the other forming part of the wall of the hypopharynx (the food tube behind the larynx, leading to the esophagus). The 102295003 |Structure of hypopharyngeal aspect of interarytenoid fold (body structure)| may be considered part of the hypopharynx, the larynx, or both. A tumor of this site is categorized as a tumor of the hypopharynx, and not the larynx, but the 105585004 |Interarytenoid fold structure (body structure)| is considered part of the larynx.

SNOMED CT does not give a Part of relationship between the hypopharyngeal aspect of the interarytenoid fold and the interarytenoid fold. This emphasizes SNOMED CT modeling based on the relationship of anatomical entities and disorders and procedures and not simply by reading term names.

Nasal turbinates
SNOMED CT differentiates between the bone underlying the nasal turbinates and the actual turbinates:

Bones underlying the turbinates,
- 118648008 |Inferior nasal turbinate bone structure (body structure)|
- 122491002 |Middle nasal turbinate bone structure (body structure)|
- 122492009 |Superior nasal turbinate bone structure (body structure)|
- 122493004 |Supreme nasal turbinate bone structure (body structure)|

Turbinates, which include bone, overlying mucous membranes, and other tissue,
- 6553002 |Inferior nasal turbinate structure (body structure)|
- 122491002 |Middle nasal turbinate bone structure (body structure)|
- 65289004 |Superior nasal turbinate structure (body structure)|
- 33415007 |Supreme nasal turbinate structure (body structure)|

The 118648008 |Inferior nasal turbinate bone structure (body structure)| is a facial bone and skull bone. And, parts of the ethmoid bone form the middle, superior, and supreme nasal conchae. This means that the bones of the middle, superior, and supreme turbinates are not bone organs.

Morphologic Abnormality Modeling
The morphologic abnormality subhierarchy is located two levels below the Body structure hierarchy with siblings Apoptosis and Structure resulting from tissue repair process:

- SNOMED CT concept
  - Body structure (body structure)
    - Body structure, altered from its original anatomical structure (morphologic abnormality)
      - Apoptosis (morphologic abnormality)
      - Morphologically abnormal structure (morphologic abnormality)
      - Structure resulting from tissue repair process (morphologic abnormality)

The concepts in the morphologic abnormality hierarchy represent abnormal body structures.

⚠️ Leave Primitive
The subhierarchy of 118956008 |Body structure, altered from its original anatomical structure (morphologic abnormality)| is to remain primitive. Authors are not to define morphologic abnormality concepts.

Content in this space
Abscess
There are two types of abscesses:
1. septic
2. sterile

Most abscesses are septic, which means that they are the result of an infection.

If a concept has a meaning (based on its FSN and text definition) that does not specify whether the abscess is sterile or septic, then the concept should not be modeled as septic; the concept’s logic definition uses 116676008 | Associated morphology (attribute) with the value 44132006 | Abscess (morphologic abnormality).

Combining morphologies

When modeling a concept requiring two role groups with the same finding site/body structure but two different morphologies (because a combined morphology does not exist), then those two morphologic abnormalities can be combined to create a new single, combined | [morphologic abnormality] | concept. Keep the newly created morphologic abnormality concept primitive as all morphologic abnormality concepts should be primitive. Limit the combination to two morphological concepts into a single concept; combining more than two morphological concepts into a single combined concept is not permitted.

Example, If 400067002 | Acantholytic epidermal nevus (disorder) | had the same Finding site of | Skin structure (body structure) | with two different morphologic abnormalities of | Epidermal nevus (morphologic abnormality) | and | Acantholysis (morphologic abnormality) |, then those two morphologic abnormality concepts can be combined to create a single, primitive, morphologic abnormality concept of | Acantholytic epidermal nevus (morphologic abnormality) |. This will prevent modeling with two relationship groups.

Instead of modeling as per this diagram in the stated view with two morphologies of the same finding site:

Model as in the stated view of this diagram with a combined morphology:
Morphologies can be combined to create a single morphologic abnormality concept where doing so creates a specialization of the morphology e.g. 55075001 |Bleeding ulcer (morphologic abnormality)|. Where morphologies are different e.g. abscess and cellulitis, they cannot be combined.

This guidance is not being applied retrospectively so the concept 707496003 |Inflammation and consolidation (morphologic abnormality)| will be an exception to this rule as it already exists as a current concept.

Combining morphologies to create a specialization is done when modeling a concept that requires two role groups with the same body structure and two morphology values. Creating a combined morphology concept enables one role group to be used.

Older 'like' content may still use the two role groups. This content would also need to use the new combined morphology value to support correct subsumption.

Neoplasm exception
Subtypes of 108369006 |Neoplasm (morphologic abnormality)| are not to be combined. These morphologies represent histological cell types that are recognized internationally by pathologists, classified by WHO, and aligned with the ICD-O classification.

Congenital anomaly
Disorders which involve congenital anomalies are defined with Occurrence (attribute) = Congenital (qualifier value), Associated morphology (attribute) = Morphologic abnormality (qualifier value) and Pathological process (attribute) = Pathological development process (qualifier value).

Therefore, congenital does not need to be represented as the Associated morphology (attribute) target value. Congenital anomaly morphology concepts usually have non-congenital parents.

Creating new morphologies
When considering the creation of a new morphological abnormality concept, consider the following:

- When the name of a potentially new morphology is the same as the disease, creation of that new morphologic abnormality concept may not be beneficial since it would not be very morphologically descriptive.
- Determine if there is benefit in creating a new morphology for a very small number of rarely used leaf nodes. If the new morphology is needed to differentiate two non-leaf concepts, that would be sufficient to create the new morphology. Otherwise, use the nearest existing morphologic abnormality.
- Including a body site in a morphological abnormality concept is forbidden unless there is a clear modeling and pathological need.

Degeneration and Degenerative abnormality
A distinction should be made between 107669003 |Degenerative abnormality (morphologic abnormality)| and 33359002 |Degeneration (morphologic abnormality)|.

- 33359002 |Degeneration (morphologic abnormality)| is a child of 107669003 |Degenerative abnormality (morphologic abnormality)|.
- 107669003 |Degenerative abnormality (morphologic abnormality)| is a grouper concept with members usually characterized by retrogressive pathologic structural changes. Diseases that are degenerative do not necessarily have 116676008 |Associated morphology (attribute)| of 33359002 |Degeneration (morphologic abnormality)|, since the word degenerative sometimes refers to loss of function rather than structural degeneration.

Examples include degeneration proper, as well as lysis, vascular sclerosis, necrosis, infarct, deposition, dystrophy, pigmentation, atrophy, and depletion.

- Morphologies under degeneration also have retrogressive structural changes, but they are not necessarily any of the above, nor are they necessarily resorption, malacia, obliteration, opacity, plaque, or postmortem change (this seems to be definition by exclusion).
- Necrosis is a degenerative abnormality, but not a degeneration. Necrosis can follow degeneration.
- Atrophy is a degenerative abnormality, but only atrophic degeneration is also a degeneration.

Modeling
33359002 |Degeneration (morphologic abnormality)| and 107669003 |Degenerative abnormality (morphologic abnormality)| should rarely, if ever, be used as the value of Associated morphology of a particular disorder; rather, a more specific subtype should be used.

Exception
It might be used as the value of Associated morphology for a broad category of degenerative disorders when the degeneration is always and necessarily structural. It is then inherited by all the subtypes, unless specialized by assigning a subtype of degeneration as the value for Associated morphology.
Fracture

Fractures should be agnostic as to whether they are pathologic or not, unless specified in the FSN or could only be caused by trauma (e.g. open fractures). Although most fractures are traumatic, there are some pathological fractures. Based on its FSN and text definition, if the word "pathological" is present, use Pathologic fracture (morphologic abnormality).

Example: Modeling Traumatic vs. Pathological Fractures

<table>
<thead>
<tr>
<th>Concept</th>
<th>Finding Site</th>
<th>Associated Morphology</th>
</tr>
</thead>
<tbody>
<tr>
<td>21351003 Fracture of phalanx of foot (disorder)</td>
<td>Bone structure of phalanx of foot</td>
<td>Fracture (morphologic abnormality)</td>
</tr>
<tr>
<td>704168008 Pathological fracture of phalanx of foot (disorder)</td>
<td>Bone structure of phalanx of foot</td>
<td>Pathologic fracture (morphologic abnormality)</td>
</tr>
</tbody>
</table>

Morphologic Abnormalities vs. Findings

Concepts from the Morphologic Abnormality hierarchy should not be used in place of concepts from the Clinical Findings hierarchy, even though they appear to refer to similar clinical situations.

For example,

- 4147007 Mass (morphologic abnormality) is not a finding, but 300848003 Mass of body structure (finding) is a finding

Morphologies are used as the values of the defining attributes of findings and procedures. Findings are used to represent the combination of a morphology and a location.

For example,

- 300923002 Cyst of scalp (disorder) represents cystic type of morphology that has the location, scalp

Many morphologies have names that could be misinterpreted as implying a process rather than a structure.

- Inflammation might mean the structural-morphologic features of inflammation, such as inflammatory cell infiltrates; or it might mean the process that causes the structural changes. Within the morphologic abnormality hierarchy, the structural interpretation is intended, not the process interpretation.

Tumor morphology

SNOMED CT accepts tumor concepts, as long as they are included in the International Classification of Diseases for Oncology (ICD-O). ICD-O has two coding systems for coding the site (topography) and the histology (morphology) of the neoplasm:

- Topographical - Anatomical site of origin or the organ system
- Morphological - Tumor cell type or histology and behavior, i.e. malignant versus benign

The topography code describes the site of origin of the neoplasms. The morphology code describes the cell type of the tumor and its biologic activity, in other words, the characteristics of the tumor itself. The morphology code, combined with the appropriate topography, expresses the complete morphological assessment as stated by the pathologist.

Specifically, there are histology types that refer to an organ by means of reference to the architecture, that is the particular histology. For example, Lymphoepithelioma-like carcinoma as enumerated by CAP (and ICD-O). The topographical reference is made to a morphology that is similar to, but distinct from, lymphoepithelium.

SCT intends to avoid adding concepts that conflate the localization of a specific tumor type in a topographic location as opposed to a neoplastic cell type that is derived from a specialized cell in an organ, e.g. adenocarcinoma vs. renal clear cell carcinoma. One is general; the other is specific to a cell type.

For more information
Clinical Finding and Disorder

Definition

| Clinical finding: normal/abnormal observations, judgments, or assessments of patients |
| Disorder: always and necessarily an abnormal clinical state |

Examples

<table>
<thead>
<tr>
<th>Clinical finding</th>
<th>Disorder</th>
</tr>
</thead>
<tbody>
<tr>
<td>Abnormal urinalysis (finding)</td>
<td>Neoplasms (disorder)</td>
</tr>
</tbody>
</table>

Clinical findings or observations are the active acquisition of subjective or objective information from a primary source. This includes information acquired from human observers, through recording of data via the use of scientific instruments or indirectly from samples taken from the source and evaluated separately.

Observations

The term "observations" should not be confused with "Observable entity", i.e. the name of something that can be observed and represents a question or assessment which can produce an answer or result (e.g. systolic blood pressure, color of iris, gender).

Context

The default context for a Clinical finding concept is:

- Present (vs. being absent)
- Subject of the record (the patient)
- Current, if not specifically stated or specified to a time in the past by an entity linked to the concept

The Clinical finding hierarchy contains the sub-hierarchy of Disorder. Concepts that are descendants of disease (disorder) are always and necessarily abnormal clinical states.

This subtype allows diseases to be subtypes of other disorders, as well as subtypes of findings.

Concepts with a semantic tag of disorder, must have a parent of Disease (disorder) and not Clinical finding (finding).

For example,

- 95617006 Neonatal cyanosis (disorder) has the parent, Disease (disorder); it is a subtype of 3415004 Cyanosis (finding).

The distinction between a disorder and a finding may be difficult to define. There are, however, distinct characteristics of each.

Disorder vs Finding

<table>
<thead>
<tr>
<th>Characteristics</th>
</tr>
</thead>
<tbody>
<tr>
<td>Disorder</td>
</tr>
<tr>
<td>Always and necessarily abnormal</td>
</tr>
<tr>
<td>Necessarily have an underlying pathological process</td>
</tr>
<tr>
<td>Have temporal persistence (may be under treatment, in remission, or inactive, even though they are still present)</td>
</tr>
<tr>
<td>May be present as a propensity for certain abnormal states to occur, even when treatment mitigates or resolves those abnormal states</td>
</tr>
<tr>
<td>Finding</td>
</tr>
<tr>
<td>May be normal (but not necessarily)</td>
</tr>
<tr>
<td>May exist only at a single point in time (e.g. a serum sodium level)</td>
</tr>
<tr>
<td>Cannot be temporally separate from the observation (one cannot observe them and say they are absent, nor can they be present when they cannot be observed)</td>
</tr>
<tr>
<td>Cannot be defined only in terms of an underlying pathological process that is present, when the observation itself is not present</td>
</tr>
</tbody>
</table>
In some cases the disease process is irrefutable, e.g. meningococcal meningitis. In others an underlying disease process is assumed based on the temporal and causal association of the disorder and its manifestation, e.g. nystagmus (disorder) is different from nystagmus present (finding). Nystagmus present (finding) may be a normal physiological response to head rotation. A person who spins around and has nystagmus present (finding), does not have nystagmus (disorder). Alternatively, a person may have nystagmus (disorder), but not nystagmus present (finding), i.e. they do not currently manifest nystagmus. Similarly, hearing loss (disorder) is different from perception of hearing loss (finding), which can be due to a number of temporary causes, such as excessive ear wax.

Clinical Finding Attributes Summary

When authoring in this domain, these are the approved attributes and allowable ranges. They are from the Human Readable Concept Model (HRCM).

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Grouped</th>
<th>Cardinality</th>
<th>In Group Cardinality</th>
<th>Range Constraint</th>
</tr>
</thead>
<tbody>
<tr>
<td>255234002</td>
<td>After (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
</tr>
<tr>
<td>116676008</td>
<td>Associated morphology (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
</tr>
<tr>
<td>47429007</td>
<td>Associated with (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..*</td>
</tr>
<tr>
<td>288556008</td>
<td>Before (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
</tr>
<tr>
<td>246075003</td>
<td>Causative agent (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
</tr>
<tr>
<td>263502005</td>
<td>Clinical course (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
</tr>
</tbody>
</table>
### Clinical Finding Defining Attributes

#### Self-grouped Attributes

Generally, the attributes before, during, after, due to, clinical course, or temporally related to are self-grouped, meaning they must not be placed in a relationship group with other attributes; each attribute must be the only attribute in a relationship group. Any rare exceptions will be documented within the individual attribute section below.

The Human Readable Concept Model (HRCM) grouped column (see the Clinical Finding Attributes Summary table on the previous page) correctly indicates that these attributes are put into a relationship group during classification because they are self-grouped.

The following defining attributes correspond to the Clinical Finding/Disorder Attributes Summary table.
After

This attribute is used to model concepts in which a clinical finding occurs after another clinical finding, procedure or event. Neither asserting nor excluding a causal relationship, it instead emphasizes a sequence of events. This attribute is self-grouped.

For example,

- 123948009 [Post-viral disorder (disorder)] occurs [After (attribute)] 34014006 [Viral disease (disorder)]

A clinical finding may start either: after a variable period of time; immediately following the resolution of its antecedent; or during the course of its antecedent but continue after the antecedent has resolved. These sequences correspond to Allen’s interval algebra relations of:

- X takes place before Y
- X meets Y
- X overlaps with Y

Associated morphology

This attribute specifies the morphologic changes seen at the tissue or cellular level that are characteristic of a disease.

(Please see Morphologic Abnormalities vs. Findings for details).

For example,

- 75694006 [Pancreatitis (disorder)] has an Associated morphology (attribute) of 409774005 | Inflammatory morphology (morphologic abnormality)]

When selecting a value for this attribute, in general, the concept should not represent a body structure combined with the morphology. There are, however, exceptions, i.e. where a morphology implies the finding site:

For example,

- Thymoma (morphologic abnormality)
- External hyperostosis (morphologic abnormality)
- Odontoma (morphologic abnormality)

Body structure should be captured in the value selected for the Finding site attribute. There are, however, exceptions.

For example,

- 70529004 [Lymphoid hyperplasia of appendix (disorder)] has [Associated morphology (attribute)] of
- 43961000 [Lymphoid hyperplasia (morphologic abnormality)] and a [Finding site (attribute)] of
- 45679000 [Appendiceal lymphoid nodule (body structure)]
Associated with

47429007 |Associated with (attribute)| represents a clinically relevant association between concepts without either asserting or excluding a causal or sequential relationship between the two. In general, avoid using the 47429007 | Associated with (attribute)| as it may be ambiguous and difficult to apply consistently.

Areas of content that use this attribute:

- Devices
- Intolerance to substances
- Concepts that group specific associations

For example,

6211002 |Polyarthritis associated with another disorder (disorder)|
Figure 2: Stated view of 6211002 |Polyarthritis associated with another disorder (disorder)| using the |Associated with (attribute)|

Before

This attribute is used to model pre-procedure complications (e.g., preoperative complication). It represents temporal associations between procedures and related disorders. This attribute is self-grouped.

Causative agent

This attribute identifies an organism, substance, physical object, physical force, and/or pharmaceutical/biological product as the direct cause of a condition. It does not include vectors, for example, a mosquito that transmits malaria.

For example,

- 4989003 |Electrical burn of skin (disorder)| has the 246075003 |Causative agent (attribute)| of 18213006 |Electricity (physical force)|

Clinical course

This attribute is used to represent both the course and onset of a disease or condition.

For example,

- 74973004 |Chronic fibrosing pancreatitis (disorder)| has a 263502005 |Clinical course (attribute)| of 90734009 |Chronic (qualifier value)|

The clinical course value is added when appropriate to the condition and thus specified in the FSN. The distinction is often necessary in those conditions that can have either an acute or a chronic course, such as bronchitis. For those conditions that have only one clinical course, i.e., diabetes is a chronic disease, a wider discussion is necessary before a decision can be made whether to assign a clinical course. Decisions on these concepts are currently made on a case-by-case basis.

Many conditions with acute (sudden) onsets also have acute (short-term) courses. Few conditions with chronic (long-term) durations require rapid versus gradual onset subtyping. Thus, there is no clear need for separating the rapidity of onset from the duration of a disease. The clinical course attribute, which combines onset and course, has been more reproducible and useful than two attributes that attempt to separate the meanings.
The word acute

The word acute has more than one meaning, and the meanings are often overlapping or unclear. It may imply rapid onset, short duration, or high severity; in some circumstances it might be used to mean all of these. For morphological concepts, acute may also imply the kind of morphology associated with the speed of onset.

For example,

- 4532008 Acute inflammation (morphologic abnormality) does not necessarily have a clinical course of sudden onset and/or short duration, but rather implies polymorphonuclear infiltration (84499006 Chronic inflammation (morphologic abnormality)) implies mononuclear cell infiltration, not necessarily a chronic course, although inflammation with a chronic course is highly correlated with a lymphocytic infiltration.

- 2704003 Acute disease (disorder) is modeled with a Clinical course (attribute) of Sudden onset AND/OR short duration (qualifier value). For clinical conditions that necessitate further specificity, the more appropriate subtypes are available.

Normally, 263502005 Clinical course (attribute) is self-grouped, meaning not grouped with other attributes. However, an exception to this rule was made for Alpha gal syndrome. Because the clinical course describes the realization and not the entire concept itself, the Clinical course (attribute) is grouped with the 719722006 Has realization (attribute).

- 788781001 Delayed allergy to red meat (finding) groups 263502005 Clinical course (attribute) of 788800008 Delayed onset (qualifier value) with 719722006 Has realization (attribute) of 769260004 Immunoglobulin E-mediated allergic process (qualifier value). Each is contained in a separate role group with a causative agent of either 226915003 Red meat (substance) or 788778006 Galactose-alpha-1,3 galactose (substance). A tick bite causes the 788779003 Allergy to galactose-alpha-1,3 galactose (finding), which in turn causes the 788781001 Delayed allergy to red meat (finding).

Due to

This attribute is used to identify a clinical finding/disorder, event, or procedure concept as the direct cause of another Clinical finding or Disorder concept. If the clinical finding merely predisposes to another disorder, rather than causing it directly, the more general Associated with (attribute) is used instead.

This attribute is self-grouped.

For example,

- 43959009 Cataract of eye due to diabetes mellitus (disorder)

During

This attribute is used to model concepts in which a clinical finding occurs during a procedure. Neither asserting nor excluding a causal relationship, it instead emphasizes a sequence of events. This attribute is self-grouped.

For example,

- 10901000087102 Hypotension during surgery (disorder) has the value Surgical procedure (procedure) for During (attribute)

Episodicity

This attribute is used to represent episodes of care provided by a physician or other healthcare provider, not episodes of disease experienced by the patient.

For example,
Asthma with 246456000 [Episodicity (attribute)] of 255217005 [First episode (qualifier value)] represents the first time the patient presents to their healthcare provider with asthma.

Finding informer
This attribute specifies the person or other entity from which the clinical finding information was obtained. It is not about the particular individual but about the category or type of informer. It is used to differentiate patient-reported symptoms from provider-determined signs. This attribute is frequently used in conjunction with 418775008 [Finding method (attribute)].

Finding method
This attribute specifies the means by which a clinical finding was determined. It includes findings that were determined by examination of the patient. Finding method is frequently used with Finding informer.

For example,

- 713071004 [Alcohol misuser in household (finding)] has the 418775008 [Finding method (attribute)] of 84100007 [History taking (procedure)]

Finding site
This attribute specifies the body site affected by a condition.

For example,

- 90708001 [Kidney disease (disorder)] has 363698007 [Finding site (attribute)] of 64033007 [Kidney structure (body structure)]

Has interpretation
This attribute refers to and designates the judgment aspect being evaluated or interpreted (e.g. presence, absence, degree, normality, abnormality, etc.). Subtypes of Environment or geographical location (environment / location) can also be used as the value in cases such as specifying a location of an incident to be reported to death and injury registries.

Interprets and Has Interpretation are grouped together in a relationship group without any other attributes.

For example,
Figure 3: Inferred view of Inadequate intake of vitamin D and vitamin D derivative (finding)

Qualifier values of |Below reference range| and |Above reference range| are preferred over values such as high/low, increased/decreased, etc. to describe Measurement finding (finding) concepts.

Interprets
This attribute refers to the entity being evaluated or interpreted, when an evaluation, interpretation, or judgment is intrinsic to the meaning of a concept.

Interprets and Has Interpretation are grouped together in a relationship group without any other attributes. Interprets may be used in a relationship group by itself without any other attributes if the value of the observable is not defined.

For example,
Figure 4: Stated view of |Decreased muscle tone (finding)|

In general, the value for the |Interprets| attribute should be from the |Observable entity| hierarchy rather than the |Procedure| hierarchy.

|Observable entity| concepts that are modeled with a |Scale type (attribute)| relationship should not be used as a value for a Clinical finding's |Interprets| relationship. The existing *vital sign* |Observable entity| concepts, e.g. Arterial blood pressure (observable entity) are exceptions to this guideline; they are permitted for use.

Be aware that SNOMED CT currently contains some concepts in the |Evaluation Procedure| hierarchy which logically belong in the |Observable entity| hierarchy. Reconciliation of the overlap between these two hierarchies will be undertaken at a future date. Discussions about the final solution for the |Observable entity| and |Evaluation Procedure| issue are ongoing. See Observable Entity vs. Evaluation procedure.

When working with the Interprets attribute, consider the values used by the supertypes and possible subtypes of your concept for this attribute. This is because the |Interprets| values must be drawn from the same hierarchy, e.g. |Observable entity| hierarchy or |Procedure| hierarchy as supertypes and subtypes, to support modeling and correct subsumption.

**Measurement finding**

For concepts in the 118245000 |Measurement finding (finding)| subhierarchy, the value for 363714003 |Interprets (attribute)| can be an Evaluation procedure, Laboratory procedure, or an Observable entity concept. In the future, the range of values may change when discussion of the relationship between evaluation procedures and observable entities concludes.

Has realization

This attribute is used to specify the process or activity that is the consequence of realization of the function.
Occurrence

This attribute refers to the specific period of life during which a condition first presents. However, conditions may persist beyond the period of life when they first present.

For example,

- 192611004 | Childhood phobic anxiety disorder (disorder) | has the 246454002 | Occurrence (attribute) of 255398004 | Childhood (qualifier value)

Pathological process

This attribute provides information about the underlying pathological process of a disorder, i.e. it describes the process that results in the structural or morphologic change.

441862004 | Infectious process (qualifier value) | and its subtype 442614005 | Parasitic process (qualifier value) | are included in the range for 370135005 | Pathological process (attribute). These are used in modeling the 40733004 | Infectious disease (disorder) subhierarchy.

For example,

- 17322007 | Disease caused by parasite (disorder) | has the 370135005 | Pathological process (attribute) of 442614005 | Parasitic process (qualifier value)

370135005 | Pathological process (attribute) must not be used for values that could overlap with 116676008 | Associated morphology (attribute).

For example,

- Inflammatory processes result in inflammation (by definition), but these disorders should be defined by their morphology, i.e. 708039003 | Inflammatory lesion (morphologic abnormality)

Disorders which involve congenital anomalies are defined with the following grouped attribute-value pairs:

- Occurrence (attribute) = congenital (qualifier value)
- Associated morphology (attribute) = << 49755003 | Morphologically abnormal structure (morphologic abnormality)>
- Pathological process (attribute) = pathological development process (qualifier value)
- Finding site = X (body structure)

Modeling

Congenital X morphology concepts should not be used. They may be used only if there is not a non-congenital supertype.
Severity

This attribute is used to subclass a Clinical finding concept according to its severity. However, this use is relative, i.e. it is incorrect to assume that the disease intensity or hazard is the same for all clinical findings to which this attribute is applied.

The severity attribute may be applied to subtypes of Clinical finding (excluding <+ Symptom severity (finding)) to represent the severity of a finding or disease.

While this attribute is useful to create subtypes of specific concepts and to differentiate the severity of a single disorder, it is not commonly used, and care must be taken when applying it. This is because:

- Severity may be interpreted in different ways, depending on the set of values available. Consider the different meaning of severity in each of the following value sets:
  - mild / moderate / severe
  - minimal / mild / moderate / severe / very severe
  - mild / mild to moderate / moderate / moderate to severe / severe / life threatening / fatal
- Severity is defined relative to the expected degree of intensity or hazard of the Clinical finding that is being qualified. For example, the common cold has a baseline intensity or hazard that is much less than a more serious disease like lupus erythematosus or pneumonia; thus, a severe cold might be considered less intense or less hazardous than mild pneumonia.
- Some disorders that are life-threatening do not ordinarily have a severity assigned to them. Cancer, for example, is not usually described as mild, moderate, or severe, but rather by stage or grade.

The Severity attribute is not applied to subtypes of 162465004 |Symptom severity (finding)| because the severity of a symptom is different to the severity of a disease. Please note this piece of guidance does not align with the MRCM but is an editorial guideline.

Modeling

Generally, 246112005 |Severity (attribute)| is not used to model concepts precoordinated in the International Release, but there are some exceptions.

A valid exception requires an internationally accepted definition that can be consistently applied and used reliably for international comparison. Even though a reference may be internationally sourced, its use may not always be uniformly applied by multiple countries. Classifications of severity that represent variation in clinical presentations and enact limitations with age ranges, sex, or pregnancy status, do not apply universally to all patients of all ages, prove problematic, and may not be generally useful.

The requestor is responsible for obtaining permission for use in SNOMED CT if required by the international body.

As an alternative to precoordination in the international release, this attribute can be used as a qualifier in postcoordination. However, beware that postcoordination of severity results in the same irreproducibility issues as pre-coordination.

Temporally related to

This attribute applies to perioperative complications in the clinical finding hierarchy. The attribute has a subhierarchy that specifies a period of time occurring before, during, or after a procedure; e.g. perioperative complications refer to complications temporally related to a surgical procedure. They include preoperative, intraoperative, and postoperative complications and are modeled with a relationship consisting of 726633004 |Temporally related to (attribute)| or an appropriate subtype. This attribute is self-grouped.

Clinical Finding and Disorder Naming Conventions

A Clinical finding/Disorder concept's fully specified name (FSN) must be specific, though the preferred term (PT) can be a more clinician-friendly, word-order variant.
The FSN must conform to a specific pattern of "Disease of x" where a specific body structure is involved. For the preferred term, end users can choose the desired description that conforms to common clinical usage.

FSN: Disease of kidney
PT: Can be either 'Kidney disease' or 'Renal disease'

The morphologic abnormality is named before naming the anatomical site.

For example,

- In 399525009 |Inflammation of ampulla of Vater (disorder)|, Inflammation is the morphologic abnormality and Ampulla of Vater is the finding site.

While the general naming convention for findings and disorders is <Morphology> of <x body structure>, there are some exceptions:

- Disorders with a well recognized name that represents the morphology; e.g. pneumonia is the well established clinical name for inflammation and consolidation of the lung
- Disorders where the meaning is not equivalent to <morphology> of <x> site convention; e.g. inflammatory bowel disease has a more specific meaning than inflammation of bowel
- Disorders which are not described by an anatomical site; e.g. metabolic disease, hereditary disease, bacterial disease

Please see documented naming patterns:

- Completed or in review - Precoordination Naming Pattern Project
- Proposed for future review - Unreviewed Naming Patterns by Hierarchy

Descriptions that include body structures

Descriptions for Clinical findings and disorders should follow the naming guidelines for Body structures if they are to be used within the Clinical finding/disorder concept. Concepts describing limbs are abundant, and the use of limb in the FSN and the synonyms of upper/lower extremity, arm/leg should be followed.

For example,

249945007 |Monoparesis of lower limb (disorder)|

Because the finding site is 61685007 |Lower limb structure (body structure)|, which follows the anatomical guidelines, the disorder concept reflects lower limb in the FSN, while using synonyms of Monoparesis of leg and Monoparesis of lower extremity.

Disorder

In the disorder hierarchy, the following naming conventions apply:

- The word disorder should be singular, so correct convention is Disorder of nose, not Disorders of nose.

Exceptions

Plurals may be used:

- As synonyms for grouper concepts, e.g. disorders or diseases
- In bilateral concepts, e.g. Disorder of bilateral eyes, Disorder of both eyes (see also Lateralized Disorder Naming Conventions)
• When the concept is a general grouping of disorders of a body system, body site, or other broad category, the word disorder is preferred over the word disease for the FSN, e.g. Disorder of reproductive system, not Disease of reproductive system. This does not apply at the leaf level.

For example,

417683006 | Sickle cell-hemoglobin C disease without crisis (disorder)

For naming conventions concerning surgical complications, sequelae, and late effects, see this section at Complication and Sequela Modeling.

Disorder X without Disorder Y
The vast majority of existing X without Y concepts originated from ICD-9 with the specific meaning of "X disorder without mention of Y disorder". As the phraseology indicates a lack of data about disorder Y as opposed to a specific exclusion, this type of concept has not been included in ICD-10, nor proposed for ICD-11, except in the case of "Traumatic brain injury without open intracranial wound".

Addition of new X without Y concepts may only be made under the following conditions:

• The request for new content must be accompanied by a rationale as to the difference between "X disorder without Y disorder" and "X disorder"
• Approval of addition by the Head of Terminology

For the most part, existing X without Y concepts will be inactivated as AMBIGUOUS with a historical MAY BE relationship to "X disorder". Exceptions to inactivation will be made on a case-by-case basis.

Requires [procedure/drug] (finding)
SNOMED international is no longer accepting new requests for concepts of the type – Requires [procedure/drug] (finding). These are administrative statuses rather than clinical findings, and this status should be represented outside of the terminology in the information model. The only exceptions relate to legacy content, and requests for subtypes of 723620004 | Requires vaccination (finding)| will continue to be accepted.

Region
If the 363698007 | Finding site (attribute) | value of a concept is a body structure with "region" in its FSN, then the description of the finding site within the clinical finding concept's FSN should also include "region".

For example, 274205003 | Burn of eye region (disorder) | has a finding site of 371398005 | Eye region structure (body structure) |

• FSN: Burn of eye region (disorder)
• PT: Burn of eye region
Allergy to substances, multiple substances

Previously, allergies caused by multiple substances were modeled by multiple causative agents suggesting that the allergy is caused by all those substances. However, when multiple substances are noted in the FSN, the intended clinical meaning is that a patient might be affected by one or more of these substances (or products containing them). To convey this meaning, these types of concepts should be modeled GClIs to represent the disjunctive meaning. e.g. 870731003 |Allergy to carbidopa and/or levodopa (finding)|

Adverse reaction caused by organisms

A description for any concept that names an organism should be consistent with the Organism hierarchy description rules.

- Use the scientific name for the organism in the FSN, e.g. Adverse reaction caused by Artemisia vulgaris pollen (disorder)
- Use the common name in the preferred term, e.g. Adverse reaction caused by mugwort pollen
- The synonym should match the FSN, e.g. Adverse reaction caused by Artemisia vulgaris pollen

Allergic and nonallergic hypersensitivity (pseudoallergic) dispositions

Allergic and nonallergic hypersensitivity (pseudoallergic) dispositions are the propensity to develop adverse allergic or nonallergic hypersensitivity (pseudoallergic) disorders. A description for any concept that names a substance or an organism should be consistent with the corresponding hierarchy description rules.

Drug allergies

Allergic and nonallergic hypersensitivity (pseudoallergic) concepts include drug allergies.
Patterns:
FSN: Allergy to X (finding)
PT: Allergy to X
For example,
  - FSN: Allergy to abacavir (finding)
  - PT: Allergy to abacavir
  - FSN: Allergy to Artemisia vulgaris pollen (finding)
  - PT: Allergy to mugwort pollen

FSN: Allergy to X and Y (finding)
PT: Allergy to X and Y
  - X and Y in alphabetical order for concepts representing multiple substances

Allergic and nonallergic hypersensitivity (pseudoallergic) disorders
These disorders represent manifestations of pathologic processes that may result in abnormal structures (e.g., allergic rhinitis).

<table>
<thead>
<tr>
<th>Disorder</th>
<th>Patterns and examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>FSN</td>
<td>Patterns:</td>
</tr>
<tr>
<td></td>
<td>• FSN: Allergic disease X (disorder)</td>
</tr>
<tr>
<td></td>
<td>• FSN: Allergic disease X (caused by Y) (disorder)</td>
</tr>
<tr>
<td></td>
<td>For example,</td>
</tr>
<tr>
<td></td>
<td>• Allergic rhinitis (disorder)</td>
</tr>
<tr>
<td></td>
<td>• Allergic conjunctivitis (disorder)</td>
</tr>
<tr>
<td></td>
<td>• Allergic rhinitis caused by grass pollen (disorder)</td>
</tr>
<tr>
<td></td>
<td>• Allergic rhinitis caused by house dust mite (disorder)</td>
</tr>
<tr>
<td>PT</td>
<td>Patterns:</td>
</tr>
<tr>
<td></td>
<td>• Allergic disease X</td>
</tr>
<tr>
<td></td>
<td>• Allergic disease X (caused by Y)</td>
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<tr>
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<td>For example,</td>
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<td></td>
<td>• Allergic rhinitis caused by grass pollen</td>
</tr>
<tr>
<td></td>
<td>• Allergic rhinitis caused by house dust mite</td>
</tr>
</tbody>
</table>

Allergic and nonallergic hypersensitivity (pseudoallergic) reactions
These disorders represent pathological processes that are defined as adverse reactions and allergic conditions with a pathological process of allergic or nonallergic hypersensitivity (pseudoallergic) process.
### Reaction Patterns and examples

<table>
<thead>
<tr>
<th>Reaction</th>
<th>Patterns and examples</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>FSN</strong></td>
<td><strong>Patterns:</strong></td>
</tr>
<tr>
<td></td>
<td>• Allergic reaction (caused by X) (disorder)</td>
</tr>
<tr>
<td></td>
<td>• Anaphylactic reaction (caused by X) (disorder)</td>
</tr>
<tr>
<td></td>
<td>• Anaphylactoid reaction (caused by X) (disorder)</td>
</tr>
<tr>
<td></td>
<td>For example,</td>
</tr>
<tr>
<td></td>
<td>• Allergic reaction caused by dye (disorder)</td>
</tr>
<tr>
<td></td>
<td>• Allergic reaction caused by pollen (disorder)</td>
</tr>
<tr>
<td><strong>PT</strong></td>
<td><strong>Patterns:</strong></td>
</tr>
<tr>
<td></td>
<td>• Allergic reaction caused by X</td>
</tr>
<tr>
<td></td>
<td>For example,</td>
</tr>
<tr>
<td></td>
<td>• Allergic reaction caused by dye</td>
</tr>
<tr>
<td></td>
<td>• Allergic reaction caused by pollen</td>
</tr>
</tbody>
</table>

### Contact hypersensitivity

Contact hypersensitivity represents a response elicited by contact of the skin or mucous membranes with a substance. The response may be immune mediated (allergic) or nonimmune (irritant) using the pathological process contact hypersensitivity process (qualifier value).

For example,
- Contact dermatitis (disorder)
- Irritant contact dermatitis (disorder)

### Intolerance to substances

An intolerance is the propensity to develop an adverse reaction to a substance. The nature of the adverse reaction can represent a variety of pathological processes but specifically excludes hypersensitivity (allergic and nonallergic hypersensitivity (pseudoallergic) reactions.

Due to the difficulty in precisely defining an intolerance pathological process, it is problematic to apply the model for hypersensitivity dispositions to defining intolerance to substance. For this reason, as well as the difficulty in associating a material agent with a disposition, substances are related to the intolerance disposition with the associated with attribute.

<table>
<thead>
<tr>
<th>Intolerance</th>
<th>Patterns and examples</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>FSN</strong></td>
<td><strong>Pattern:</strong></td>
</tr>
<tr>
<td></td>
<td>• Intolerance to X (finding)</td>
</tr>
<tr>
<td></td>
<td>Example,</td>
</tr>
<tr>
<td></td>
<td>• Intolerance to milk (finding)</td>
</tr>
<tr>
<td><strong>PT</strong></td>
<td><strong>Pattern:</strong></td>
</tr>
<tr>
<td></td>
<td>• Intolerance to X</td>
</tr>
<tr>
<td></td>
<td>Example,</td>
</tr>
<tr>
<td></td>
<td>• Intolerance to milk</td>
</tr>
</tbody>
</table>

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Inadequate and excessive intake of energy and nutrients

Identification of findings of inadequate or excessive intake of nutrients inconsistent with nutrient requirements and established reference standards includes nutrients with a variety of forms where applicable.

For example, 870465001 | Excessive intake of vitamin A and vitamin A derivative (finding)
FSN: Excessive intake of vitamin A and vitamin A derivative (finding)
PT: Excessive intake of vitamin A and vitamin A derivative

Laterized Disorder Naming Conventions

For more information
See also Anatomical Structure Naming Conventions section Naming Convention for Digits of Hand and Foot and Laterality section Laterality

Right, left disorder concepts
When creating a lateralized disorder concept, two concepts should be created:

1. concept for the left side
2. concept for the right side

Descriptions

- FSN: <morphologic abnormality> of right/left <body structure> (disorder)
- PT: Right/left <disorder>

For example, 1089071000119109 | Inflammation of left mastoid (disorder)

- FSN: Inflammation of left mastoid (disorder)
- PT: Left mastoiditis
Figure 1: Stated view of Inflammation of left mastoid (disorder)

Where the disorder is left/right of a specific anatomical site, and the preferred term naming pattern of Right/ left <disorder> causes a combination that does not sound like natural flowing English, the guidance above can be circumvented. See the section Naming Convention for Digits of Hand and Foot and Laterality section Laterality.

For example,

Left interphalangeal thumb joint open traumatic dislocation should follow naming guidance of Open traumatic dislocation of interphalangeal joint of left thumb.

Bilateral disorder concepts

Descriptions

- FSN: <Morphologic abnormality) of bilateral <body structure> (disorder)
- PT: Bilateral <disorder>
- SYN: <Disorder> of bilateral <body structure>
- SYN: <Disorder> of both <body structure>

For example, 1084011000119100 |Inflammation of bilateral mastoids (disorder)|

- FSN: Inflammation of bilateral mastoids (disorder)
- PT: Bilateral mastoiditis
- SYN: Inflammation of bilateral mastoids
- SYN: Inflammation of both mastoids

Modeling of bilateral disorders

Bilateral disorders should be modeled using two relationship groups, one for each lateralized body structure.
Figure 2: Stated view of Inflammation of bilateral mastoids (disorder) with a role group for each side

⚠️ **Structure, Structure of**

Lateraled disorder concepts should not include the words *structure* or *structure of*.

For example,

- For the body structure concept, 266005 | Structure of lower lobe of right lung (body structure), a disorder concept with this body structure is 724056005 | Malignant neoplasm of lower lobe of right lung (disorder).
- For the body structure concept, 266005 | Structure of lower lobe of right lung (body structure), a procedure with this body structure is 726425007 | Lobectomy of lower lobe of right lung (procedure).
Disorder Modeling

A disorder is always and necessarily an abnormal clinical state.

Disorder modeling information is as follows:

Specific Disorder Modeling

Acquired abnormality of congenital anomaly

For those concepts that describe a congenital anomaly that has been repaired and subsequently acquired an abnormality, follow the naming convention of |Acquired abnormality of X following repair of congenital X (disorder)|.

For example,
- 871598001 |Acquired abnormality of common arterial trunk following repair of truncus arteriosus (disorder)|

Adverse reaction to X vaccine

Overview

The following modeling and terming guidelines apply to concepts in the International Release.

Modeling (stated view)

"Adverse reaction to X vaccine" concepts shall be modeled using the proximal primitive modeling pattern. Due to the small number of concepts (n<50), no template will be created. The "Adverse reaction to substance" template can be consulted for generalized modeling guidance.

<table>
<thead>
<tr>
<th>Attribute: Causative agent</th>
</tr>
</thead>
<tbody>
<tr>
<td>Range: &lt;&lt;787859002</td>
</tr>
<tr>
<td>Exceptions: none identified</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attribute: Causative agent</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cardinality: 1..1</td>
</tr>
<tr>
<td>Adverse reaction to X vaccine concepts should have one and only one</td>
</tr>
<tr>
<td>Concepts representing &quot;vaccine product containing only&quot; should not be used in modeling Adverse reaction to X vaccine concepts.</td>
</tr>
<tr>
<td>Exceptions: none identified</td>
</tr>
</tbody>
</table>

GCI

Not applicable

Terming Guidelines
Use the following pattern for the FSN; align terming and case sensitivity with the FSN for the concept that represents the vaccine product that is the cause of the adverse reaction.

- Adverse reaction to component of `<Causative agent FSN>` (disorder)

Example:

- Adverse reaction to component of vaccine product containing Hepatitis A virus antigen (disorder)
- Adverse reaction to component of vaccine product containing Streptococcus pneumoniae antigen (disorder)
- Adverse reaction to component of vaccine product containing only Clostridium tetani and Corynebacterium diphtheriae antigens (disorder)
- Adverse reaction to component of vaccine product containing Measles morbillivirus and Mumps orthorubulavirus and Rubella virus antigens (disorder)

Use the following pattern for the PT; align terming and case significance with the PT for the disorder that is the target of the vaccine. For multiple ingredient vaccine products, the disorders must be listed in alphabetical order and separated by the word "and".

- Adverse reaction to `<disorder>` vaccine
- Adverse reaction to `<disorder>` and `<disorder>` vaccine
- Adverse reaction to `<disorder>` and `<disorder>` and `<disorder>` vaccine

Example:

- Adverse reaction to hepatitis A vaccine
- Adverse reaction to pneumococcal vaccine
- Adverse reaction to diphtheria and tetanus vaccine
- Adverse reaction to measles and mumps and rubella vaccine

**Synonyms**

- A synonym corresponding to the FSN is required.

- Synonyms beginning with the disorder that is the target of the vaccine are allowed. For multiple ingredient vaccine products, the disorders must be listed in alphabetical order and separated by the word "and". Note that these are not true synonyms; they may be updated and identified as "near-synonym" descriptions when that functionality becomes available although that would also potentially require updating the PT.

Example:

- Hepatitis A vaccine adverse reaction
- Pneumococcal vaccine adverse reaction
- Diphtheria and tetanus vaccine adverse reaction
- Measles and mumps and rubella vaccine adverse reaction

**Exemplars**

The following illustrates the stated and inferred view for top level grouper 293104008 |Adverse reaction to vaccine product (disorder)|:
The following illustrates the stated view for top level grouper 219075006 |Adverse reaction to vaccine product containing bacteria antigen (disorder)|:

The following illustrates the inferred view for top level grouper 219075006 |Adverse reaction to vaccine product containing bacteria antigen (disorder)|:

The following illustrates the stated view for single ingredient vaccine 293126009 |Adverse reaction to vaccine product containing Hepatitis A virus antigen (disorder)|:

The following illustrates the inferred view for single ingredient vaccine 293126009 |Adverse reaction to vaccine product containing Hepatitis A virus antigen (disorder)|:
The following illustrates the stated view for multiple ingredient vaccine 293125008 |Adverse reaction to vaccine product containing Measles morbillivirus and Mumps orthorubulavirus and Rubella virus antigens (disorder)|:

The following illustrates the inferred view for multiple ingredient vaccine 293125008 |Adverse reaction to vaccine product containing Measles morbillivirus and Mumps orthorubulavirus and Rubella virus antigens (disorder)|:

**Allergy to X vaccine**

**Overview**

The following modeling and terming guidelines apply to concepts in the International Release.

**Modeling (stated view)**

"Allergy to X vaccine" concepts shall be modeled using the proximal primitive modeling pattern. Due to the small number of concepts (n<25), no template will be created. The "Allergy to substance" template can be consulted for generalized modeling guidance.

<table>
<thead>
<tr>
<th>Single or multiple ingredient vaccine</th>
</tr>
</thead>
<tbody>
<tr>
<td>Stated parent concept</td>
</tr>
<tr>
<td>420134006</td>
</tr>
<tr>
<td>Exceptions: none identified</td>
</tr>
<tr>
<td>Semantic tag</td>
</tr>
<tr>
<td>(finding)</td>
</tr>
<tr>
<td>Definition status</td>
</tr>
<tr>
<td>90000000000000074008</td>
</tr>
<tr>
<td>Note: Because 'Allergy to X vaccine' represents the propensity to an allergic reaction to any component (including excipients) of a vaccine rather than the modeled active ingredient(s), these concepts cannot be sufficiently defined. As a result, there will not be subsumption between &quot;Allergy to X vaccine&quot; concepts.</td>
</tr>
<tr>
<td>Exceptions: Grouper concept 863903001</td>
</tr>
</tbody>
</table>

| Attribute: |
| Has realization |
| Attribute value = 472964009 |Allergic process [qualifier value]| |
| Exceptions: none identified |
### Attribute: Causative agent

- **Range:** 787859002 [Vaccine product (medicinal product)]
  - Exceptions: none identified
- **Cardinality:** 1..1
  - Allergy to X vaccine concepts should have one and only one [Causative agent] attribute.
  - Concepts representing "vaccine product containing only" should not be used in modeling Allergy to X vaccine concepts.
  - Exceptions: none identified

### GCI
Not applicable

### Terming Guidelines

#### FSN

Use the following pattern for the FSN; align terming and case sensitivity with the FSN for the concept that represents the vaccine product that is the cause of the allergy.

- **Allergy to component of** `<Causative agent FSN>` (finding)
  
  Example:
  
  - Allergy to component of vaccine product containing Hepatitis A virus antigen (finding)
  - Allergy to component of vaccine product containing Streptococcus pneumoniae antigen (finding)
  - Allergy to component of vaccine product containing Clostridium tetani and Corynebacterium diphtheriae antigens (finding)
  - Allergy to component of vaccine product containing Measles morbillivirus and Mumps orthorubulavirus and Rubella virus antigens (finding)

#### Preferred Term

Use the following pattern for the PT; align terming and case significance with the PT for the disorder that is the target of the vaccine. For multiple ingredient vaccine products, the disorders must be listed in alphabetical order and separated by the word "and".

- Allergy to `<disorder>` vaccine
- Allergy to `<disorder>` and `<disorder>` vaccine
- Allergy to `<disorder>` and `<disorder>` and `<disorder>` vaccine

Example:

- Allergy to Hepatitis A vaccine
- Allergy to pneumococcal vaccine
- Allergy to diphtheria and tetanus vaccine
- Allergy to measles and mumps and rubella vaccine

**NOTE:** For national extensions modeling using "vaccine containing only" product concepts, these disorder-based descriptions will need to reflect "only" to eliminate duplicate descriptions.
Synonyms

- A synonym corresponding to the FSN is required.
- Synonyms beginning with the disorder that is the target of the vaccine are allowed. For multiple ingredient vaccine products, the disorders must be listed in alphabetical order and separated by the word "and". Note that these are not true synonyms; they may be updated and identified as "near-synonym" descriptions when that functionality becomes available although that would also potentially require updating the PT.

Example:

- Hepatitis A vaccine allergy
- Pneumococcal vaccine allergy
- Diphtheria and tetanus vaccine allergy
- Measles and mumps and rubella vaccine allergy

NOTE: For national extensions modeling using "vaccine containing only" product concepts, these disorder-based descriptions will need to reflect "only" to eliminate duplicate descriptions.

Exemplars

The following illustrates the stated view for top level grouper 863903001 |Allergy to component of vaccine product (finding)|:

![Diagram of stated view for 863903001]

The following illustrates the inferred view for top level grouper 863903001 |Allergy to component of vaccine product (finding)|:

![Diagram of inferred view for 863903001]

The following illustrates the stated view for 294663006 |Allergy to component of vaccine product containing Hepatitis A virus antigen (finding)|:
The following illustrates the inferred view for 294663006 |Allergy to component of vaccine product containing Hepatitis A virus antigen (finding)|:

![Diagram](image1)

The following illustrates the stated view for 294662001 |Allergy to component of vaccine product containing Measles morbillivirus and Mumps orthorubulavirus and Rubella virus antigens (finding)|:

![Diagram](image2)

The following illustrates the inferred view for 294662001 |Allergy to component of vaccine product containing Measles morbillivirus and Mumps orthorubulavirus and Rubella virus antigens (finding)|:

![Diagram](image3)

**Arrhythmia**

Cardiologists noted confusion in the placement of *Conduction disorder of the heart (disorder)* as a broad grouping that subsumed arrhythmias and heart blocks. In common usage *arrhythmia* refers to a broad set of conditions that include conduction disorders, under which are heart blocks. The concept Cardiac arrhythmia (disorder) is a parent
of Conduction disorder of the heart (disorder), and the active referent of the inactive concepts named dysrhythmia or arrhythmia.

For example,

- Arrhythmias, like |72654001| Supraventricular arrhythmia (disorder), are under |698247007| Cardiac arrhythmia (disorder).

Conduction disorders include heart block, AV block, bundle branch block, conduction delay, and conduction defect, like |418341009| Atroventricular conduction disorder (disorder). Other arrhythmias were moved from under |44808001| Conduction disorder of the heart (disorder) and placed under |698247007| Cardiac arrhythmia (disorder).

Bacterial disorders with organism or toxin
In modeling some bacterial disorders, there will be situations where either the organism or the toxin (substance), or both values, are required for the causative agent attribute. The decision is often determined by whether or not the bacteria are considered endotoxins or exotoxins. The most common exotoxins are:

- Botulinum Toxin
- Enterotoxin
- Cholera Toxin
- Diphtheria Toxin
- Tetanospasmin

Exotoxins are more lethal in comparison to endotoxins, but there are vaccines against many exotoxins whereas there are no vaccines against endotoxins. There can be instances where an infection is present but the disease-causing toxins are not; in this case, model the concept only with the organism and not the toxin substance.

Example,

|276202003| Infection caused by Clostridium tetani (disorder) is modeled with a causative agent of |30917009| Clostridium tetani (organism) only.

In the situation where a disease is caused by both the infection and the associated toxin, model with both the causative agent and the toxin substance.

Example,

|76902006| Tetanus (disorder) is modeled with a causative agent of |30917009| Clostridium tetani (organism) as well as |26159005| Clostridium tetani toxin (substance).

Combining Morphologic Abnormalities
When modeling a concept requiring two role groups with the same body structure but two different morphologies (because a combined morphology does not exist), then those morphologic abnormalities can be combined to create a single |(morphologic abnormality)| concept. Keep the newly-created morphologic abnormality concept primitive as all morphologic abnormality concepts are primitive.

<table>
<thead>
<tr>
<th>Example: Modeling</th>
</tr>
</thead>
<tbody>
<tr>
<td>Concept</td>
</tr>
<tr>
<td>400067002</td>
</tr>
</tbody>
</table>

Another example is |1076491000119102| Nontraumatic complete rupture of muscle or tendon structure of rotator cuff of left shoulder (disorder).

If this disorder had the same finding site of |Structure of rotator cuff of left shoulder (body structure)| with two different morphologic abnormalities of |Nontraumatic rupture| and |Complete rupture|, then those two morphologic abnormality concepts can be combined to create a single, primitive, morphologic abnormality concept of |Nontraumatic complete rupture (morphologic abnormality)|. This will prevent modeling with two relationship groups.
Instead of modeling as in this stated view:

Model as shown in this stated view:
The concept 66091009 | Congenital disease (disorder) | means present at birth. Though the word congenital may be applied to genetic disorders, the term genetic is preferred for those disorders.

The logical definition of a congenital disorder must include:

- Occurrence = Congenital (qualifier value).
- It may also include:
  - Finding site = X (body structure)
  - Associated morphology = X (morphologic abnormality)
  - Pathological process = Pathological development process (qualifier value)

All of these defining relationships should be grouped to indicate that the abnormal morphology occurs at the finding site, results from a pathological development process, and is present at birth. Where a morphologic abnormality occurs at more than one finding site, or one body structure has multiple morphologic abnormalities, multiple relationship groups should be created and the pathological process and occurrence relationships included in each relationship group.

The following guidelines apply:

A disorder with the word congenital in the FSN should classify under 66091009 | Congenital disease (disorder).

Congenital X (morphologic abnormality) concepts are being inactivated hence Congenital anomaly grouper concepts, such as 9904008 | Congenital anomaly of cardiovascular system (disorder), should be modeled with an Associated morphology (attribute) of 49755003 | Morphologically abnormal structure (morphologic abnormality) and a Pathological process relationship.

Whether creating new or revising existing concepts, only use Congenital X (morphologic abnormality) concepts if no non-congenital supertype of that morphologic abnormality is active.

- For example, use 399898009 | Misalignment (morphologic abnormality) | not 102283003 | Congenital misalignment (morphologic abnormality).

### Neonatal period

According to the American Medical Association, the periods of life in the postnatal period include all periods after birth including the neonatal or immediate postpartum period. It may be challenging to differentiate a congenital disorder from a neonatal disorder. A condition may be present at birth, i.e. congenital; however, clinical manifestations may take longer to appear, i.e. during the neonatal period (e.g. 14333004 | Alloimmune neonatal neutropenia (disorder)).

Congenital versus acquired

While some disorders are only congenital or only acquired, some disorders may be either congenital or acquired. The acquired form should only exist when there is a need to differentiate from the congenital form. Do not model a disorder as acquired if a congenital variant does not exist.

Congenital disorders are modeled using 246454002 | Occurrence (attribute) of 255399007 | Congenital (qualifier value). If the FSN does not include congenital, it should not be modeled as congenital. The precise meaning of the FSN should be followed (e.g. many hereditary disorders have congenital appearances).

For example,

- 33534005 | Congenital bowing of femur (disorder) | is modeled with 246454002 | Occurrence (attribute) of 255399007 | Congenital (qualifier value).
Acquired disorders are those that originate and manifest after birth. The disorders are associated with a period of life, as opposed to a specific process or structure. All diseases (disorders) that occur after birth are considered acquired.

Generally, concepts that explicitly state acquired in the FSN or in a synonym should be modeled with Occurrence = 767023003 | Period of life beginning after birth and ending before death (qualifier value)|.

For example, 240253004 | Acquired abduction deformity of foot (disorder) | has acquired in the FSN and is modeled with Occurrence = 767023003 | Period of life beginning after birth and ending before death (qualifier value)|.

Remodeling Acquired Disorders

When revising acquired disorders, remove any acquired morphologies and replace with general parent morphologies, e.g. replace 127560004 | Acquired deformity (morphologic abnormality) | with 6081001 | Deformity (morphologic abnormality) |. Then add Occurrence attribute with a value of 767023003 | Period of life beginning after birth and ending before death (qualifier value)|. One of its children may also be used if the FSN states the period of life, such as Childhood or Adulthood.

Congenital absence
Congenital absence can represent at least three different classes of absence
1. Total developmental absence of the affected organ/structure
2. Partial absence of the affected organ/structure
3. In utero amputation of all or part of the affected organ/structure

In order to conform to the intended meaning of the FSNs as described by the original source, the following modeling patterns are proposed for congenital absence terms:

**Congenital absence of X**
- Associated morphology (attribute) = 418560003 |Absence (morphologic abnormality)|
- Occurrence (attribute) = 255399007 |Congenital (qualifier value)|
- Finding site (attribute) = Structure of X

**Aplasia or Partial absence of X**
- Associated morphology (attribute) = 45486003 |Aplasia (morphologic abnormality)| or 890175002 |Transverse deficiency (morphologic abnormality)|
- Occurrence (attribute) = 255399007 |Congenital (qualifier value)|
- Finding site (attribute) = Part of X
- Pathological process (attribute) = Pathological developmental process (qualifier value)

**Agenesis of X or Complete absence of X**
- Associated morphology (attribute) = 782173000 |Agenesis (morphologic abnormality)|
- Occurrence (attribute) = 255399007 |Congenital (qualifier value)|
- Finding site (attribute) = Entire X
- Pathological process (attribute) = Pathological developmental process (qualifier value)

See also relative sections:
- Acquired abnormality of congenital anomaly
- Malformation, deformation, anomaly

**Death**
*Death* is an event, not a disorder.

**Sudden cardiac death**
*Sudden cardiac death* is a term used in clinical practice. It refers to an arrhythmia that results in sudden loss of cardiac function which, if not quickly reversed, will lead to *actual* death. The FSN Sudden cardiac death (disorder) is modeled as a subtype of 127337006 |Acute heart disease (disorder)|. It should not be classified as *death*. Individuals with sudden cardiac death have not necessarily been declared *dead* and are frequently revived. It is regarded as a *subtype of cardiac dysrhythmia*.

**Fetal findings and disorders**
When modeling a fetal finding or fetal disorder, the [Finding site (attribute)] should not be a *fetal* body structure unless it is unique to the fetal period, such as [Umbilical cord structure (body structure)]. The fetal aspect is captured using the [Occurrence (attribute)] = [Fetal period (qualifier value)], as in [Fetal tachycardia (disorder)].

Note: This guideline has not been applied to fetal procedures at this point in time.

**Genetic, developmental, congenital, and physical origin**
The following figure shows the structure of genetic, developmental, and congenital categories, along with non-genetic, non-developmental, and postnatal categories. A dimension, called *extrinsic physical force*, is included to distinguish *deformations* from *malformations*. The sections of the diagram represent categories formed from the combination of the dimensions, each which represents the answer to one of the following questions:

- Is it genetic or not?
- Is it developmental or not?
- Is it present at birth or not?
- Is it due to an extrinsic physical force or not?
Figure 1: The relationships of genetic, congenital, developmental, and acquired disorders

Explanation of Figure

The sections with diagonal hashed lines represent combination categories that do not occur. For example, there are no genetic disorders that are due to an extrinsic physical force. Likewise, there are no congenital disorders that are considered non-developmental.
The sections with blue crossing lines represent congenital malformations; they may be either genetic or non-genetic.

For example, congenital infectious malformations

The red circle represents congenital genetic malformations.

The blue sections represent acquired, i.e. disorders that are non-genetic and not present at birth.

For example, Vitamin D deficiency (rickets) in children is a non-genetic, non-congenital, developmental malformation.

The white sections represent genetic congenital or genetic postnatal disorders.

For example, Huntington’s disease is a genetic disease that is neither congenital nor developmental. The gene defect is present at birth, but the disease does not manifest until adulthood.

Arrows leading from the sections point to examples of disorders for the category.

Developmental

Developmental is a useful label for disorders that affect developing structures or functions that may occur during pre- or postnatally. They may be present at birth or develop later.

Familial

The term familial may also be ambiguous when used for broad categories. It may mean that the disorder is found in higher proportions in the immediate or extended family compared to other groups. Or, it may mean there is a possibility of a disease being inherited. It may be used; however, it may require clarification of meaning from the requestor. It should not be used as a synonym for genetic.

Hereditary

It may be a challenge to classify a condition as a Hereditary disease (disorder). Hereditary requires case-by-case definition; it cannot be applied to broad categories. Nevertheless, the names by which many diseases are known include the term, and it is permitted, as long as it does not introduce ambiguity.

Hematologic and lymphatic conditions

Hematologic, lymphatic

There is more than one meaning of hematologic. A definition based on hematological system structure includes hematopoietic and lymphoid structures (including bone marrow, spleen, thymus, lymph nodes, etc), as well as the cellular components of blood. Hematologic neoplasms clearly fit this definition.

A definition based on clinical usage by hematologists is broader. Disorders of hemostasis and thrombosis are often managed by hematologists, but these do not have a common structural overlap with the lymphoid and hematopoietic systems (with the exception of platelets and megakaryocytes). For clarity, hematologic disorder is a navigational concept that is used to define a reference set that includes disorders of blood and blood forming organs, as well as disorders of hemostasis and thrombosis, depending on what is intended.

Hematologic disorders, lymphoid and myeloid neoplasms

Hematologic disorders may refer to disorders of: hematopoietic cell origin; blood forming organs (bone marrow, lymph nodes, spleen, thymus, and other lymph tissues); cellular components of blood; or function of hemostatic and thrombotic systems.

Diseases of the blood forming organs (bone marrow, lymph nodes, etc.) can be defined by any one or a combination of the following:

The morphology (neoplastic diseases, at a minimum, include those morphologies covered by neoplasms in the International Classification of Diseases for Oncology, ICD-O).

For example,

- 118599009 |Hodgkin's disease (disorder)| has 128930002 |Hodgkin lymphoma - category (morphologic abnormality)|. The body site involved (especially specific lymph node groups or skin sites).
For example,

- 400122007 | Primary cutaneous T-cell lymphoma (disorder) | has Finding site, skin structure (body structure)

For some disorders, like T-cell lymphomas, and plasma cell and immunosecretory disorders, it is important to distinguish those defined by morphological appearance, site, or manifestation.

T-cell lymphomas can be subcategorized according to the primary site, a lymph node, the skin, or other extranodal site. This means that a site of lymphoid structure cannot be the defining characteristic of the parent concept T-cell lymphoma. Its defining attribute should be morphology alone.

Plasma cell and immunosecretory disorders (e.g. monoclonal gammopathy, heavy chain disease, Waldenstrom’s macroglobulinemia) are defined by their manifestations, i.e. the type of monoclonal protein they secrete. Others (e.g. myeloma, plasmacytoma) are defined by their morphology, regardless of whether or not they are secretory.

Immunosecretory disorders may have a morphology of plasma cell neoplasm, even though no mass has been identified and the monoclonal protein may be the only evidence that there is a clonal neoplasm.

In general, lymphoid and myeloid neoplasms can be modeled with their morphologies, but without a site. Leukemias and myelodysplastic syndromes are modeled with Finding Site, bone marrow structure (body structure).

Coagulation, hemostasis, thrombosis
There is more than one meaning of coagulation. A broad meaning, to stop bleeding, is better described as hemostasis. A more narrow definition, limited to the formation of the fibrin clot, might exclude certain components of hemostasis (e.g. the ability to stop hemorrhage through the actions of blood vessels, collagen, endothelial cells, and platelets, in the absence of clotting). Individuals with congenital fibrinogen deficiency cannot form fibrin clots, yet their bodies are able to stop bleeding. Therefore, coagulation disorders are kinds of hemostatic disorders.

Hernia
Hernias involve two body structures, one is the hernial opening and the other is the herniated structure. When modeling hernias, use two role groups to represent the body structures and the respective associated morphology for each site. If the herniated structure is not explicit, use the supertype concept for the finding site.

For example,

The concept 50063009 | Femoral hernia (disorder) | is modeled with Finding site = 818983003 | Structure of abdominopelvic cavity and/or content of abdominopelvic cavity and/or anterior abdominal wall (body structure) | to represent the herniated structure.

![Figure 1: Stated view of Femoral hernia (disorder)](image)

Iatrogenic
Adding further concepts to the iatrogenic disorder hierarchy is discouraged. Concepts must have iatrogenic in the FSN to be modeled with an IS_A relationship to 12456005 | Iatrogenic disorder (disorder) |. An iatrogenic disorder should remain as a primitive concept if dependent only upon parent relationships to describe the
disorder. In cases where the modeling is explicit, e.g. 202762009 [iatrogenic cervical spinal stenosis (disorder)], the concept can be defined.

For example,

![Diagram of Iatrogenic Cervical Spinal Stenosis](image)

Figure 1: Stated view of [iatrogenic cervical spinal stenosis (disorder)] using IS_A 12456005 [iatrogenic disorder (disorder)]

Immune function disorders
Hypersensitivity
473010000 [Hypersensitivity condition (finding)] is a primitive concept. It subsumes 473011001 [Allergic condition (finding)] and 609405001 [Non-allergic hypersensitivity condition (finding)].

473010000 [Hypersensitivity condition (finding)] is a direct descendant of 404684003 [Clinical finding (finding)].

473011001 [Allergic condition (finding)] and 609405001 [Non-allergic hypersensitivity condition (finding)] are both primitive concepts. Each has three main subhierarchies representing:

- Diseases/disorders: abnormal structures
- Processes: allergic and nonallergic hypersensitivity (pseudoallergic) reactions
- Dispositions: propensities to develop allergic and nonallergic hypersensitivity (pseudoallergic) reactions; they do not have pathophysiologic manifestations prior to allergic and nonallergic hypersensitivity (pseudoallergic) processes, i.e. reactions

Diseases/disorders and reactions, but not dispositions, are defined by underlying pathological processes.

Pathological process (qualifier value) hierarchy

In order to fully describe the full range of hypersensitivity responses, there are qualifier values in the Pathological process (qualifier value) hierarchy. (See also Qualifier Value page).

Allergic reaction

Allergic reaction (disorder) has a Causative agent (attribute) of Substance (substance) or its subtypes. This attribute-value pair is grouped with another attribute-value pair of Pathological process (attribute) and Allergic process (qualifier value).
Allergic process (qualifier value) is a subtype of Abnormal immune process (qualifier value) which means allergic disorders, as well as autoimmune disorders, classify as types of disorders of immune function. Disorder of immune function (disorder) modeling with Abnormal immune process (qualifier value) allows allergic and autoimmune disorders to correctly classify as subtypes of Disorder of immune function (disorder).

Allergic and nonallergic hypersensitivity (pseudoallergic) disease
Allergic and nonallergic hypersensitivity (pseudoallergic) diseases represent manifestations of pathologic processes that result in abnormal structures. Modeling an allergic and nonallergic hypersensitivity (pseudoallergic) disease includes the following relationship group:

IS A: Disease (disorder)
Associated morphology (attribute): subtype of Morphologically abnormal structure (morphologic abnormality) representing the abnormal structure
Finding site (attribute): subtype of Anatomical or acquired body structure (body structure) representing the abnormal structure
Pathological process: Hypersensitivity process (qualifier value) or one of its descendants
Causative agent (attribute): Substance (substance) or one of its descendants, if known

For example,
Allergic and nonallergic hypersensitivity (pseudoallergic) disposition

Allergic and nonallergic hypersensitivity (pseudoallergic) dispositions are propensities to develop allergic and nonallergic hypersensitivity (pseudoallergic) reactions; they do not have pathophysiologic manifestations prior to reactions. They are considered clinical findings, not disorders. This further distinguishes them from allergic and nonallergic hypersensitivity (pseudoallergic) reactions.

Allergy to X (finding) will have the following modeling:

- **IS A:** Propensity to adverse reaction (finding)
- **Role group of:**
- **Has realization (attribute):** Allergic process (qualifier value)
- **Causative agent (attribute):** subtype of Substance (substance)

For example,
For example,

Nonallergic hypersensitivity (pseudoallergic) reaction
Nonallergic hypersensitivity (pseudoallergic) reactions are adverse reactions; they are defined by an underlying pathological process.

For example,
Intolerance to substance

An intolerance is the propensity to develop an adverse reaction to a substance. The adverse reaction may be associated with various pathological processes, but specifically excludes hypersensitivity reactions.

It may be difficult to define the pathological process and to associate the substance with the propensity to develop a reaction. Consequently, 47429007 | Associated with (attribute) | is used to model intolerance to substances.

Infectious vs. inflammatory

Disorders with the suffix "-itis" (e.g. cystitis, prostatitis, tonsillitis, appendicitis) are often infectious as well as inflammatory in nature. For inflammatory conditions whose FSNs specify an infective cause, the modeling should include:

- [Causative agent (attribute)] with the specified organism
- [Pathological process (attribute)] with the type of infectious process
An infectious cause should neither be assumed nor modeled when the FSN does not specify it. For inflammatory conditions whose FSNs do not specify an infective cause, the modeling should exclude a Causative agent and Pathological process and should include only:

- Associated morphology (attribute) of Inflammatory morphology or subtype
- Finding site (attribute) with a body structure when known

Example of inflammatory and infectious disorder:
441551009 |Inflammation of larynx caused by virus (disorder)| (synonym, Viral laryngitis) includes a |Causative agent (attribute)| of Virus (organism) and a |Pathological process (attribute)| of |Infectious process (qualifier value)|.

Example of inflammatory disorder not specified as infectious:
446292002 |Necrotizing inflammation of lymph node (disorder)| (synonym, Necrotizing lymphadenitis) does not specify an infective cause, so it is neither modeled with Causative agent nor Pathological process. The model contains an |Associated morphology (attribute)| and a |Finding site (attribute)|.

Ischemia
Ischemic disorder
Ischemic disorders are defined by a morphology of ischemic structural change. This need not be permanent, but it is assumed that all ischemia results in some structural alterations at the molecular level.

Ischemic heart disease
Ischemic heart disease includes myocardial infarction, myocardial ischemia (without infarction), angina, and other disorders of the heart that have ischemic structural change (reversible or non-reversible) as a defining characteristic.

Coronary arteriosclerosis can, of course, be present without causing ischemia, so coronary arteriosclerosis is not a subtype of ischemic heart disease.

Likewise, there are causes of myocardial ischemia and infarction other than coronary arteriosclerosis, so ischemic heart disease is not a subtype of coronary arteriosclerosis.

**Lesion**
The word lesion can be used to refer to both structural and functional abnormalities. If a disorder (or procedure) refers to a lesion in a way that makes it clear that it is a generic term for a structural abnormality, then the correct modeling approach is to use 116676008 | Associated morphology (attribute) | 49755003 | Morphologically abnormal structure (morphologic abnormality) | (For procedures, use 405816004 | Procedure morphology (attribute)).

Functional lesions should not be modeled using values from the Morphologically abnormal structure hierarchy.

**Malformation, deformation, anomaly**
The word anomaly is, by itself, ambiguous. It may mean: any abnormality including non-structural ones; malformation; both malformation and deformation. Concepts with the word anomaly must be evaluated for ambiguity.

A deformation is a structural abnormality that is due to an extrinsic physical force. Newly created concepts representing a deformity should be considered disorders.

A malformation is a structural abnormality that results from intrinsically disordered development.

For example,

- *Congenital anomaly of <x structure>* is definitely structural but is not the same as *congenital malformation* (structural abnormality due to intrinsically disordered development present at birth). Therefore, it can be regarded as having the more general meaning of *structural abnormality present at birth*.

See also relative section:
Genetic, developmental, congenital, and physical origin

Acquired abnormality of congenital anomaly

Congenital

Mental health
Dependence-related concepts which express the current existence of abuse are acceptable.

For example,

- 191816009 | Drug dependence (disorder) |

Dependence-related concepts which express the pattern as either continuous or episodic are not acceptable.

Unacceptable patterns,

- X with single episode
- X with multiple episodes
- Current episode of X
- First episode of X
- X with continuous pattern

Unacceptable legacy concepts,

- Drug abuse, continuous (disorder)
- Episodic drug abuse (disorder)

Concepts describing full or partial remission are acceptable but not the phase of the remission. The patterns are:
• X in full remission
• X in partial remission

For example,
• 46244001 | Recurrent major depression in full remission (disorder) |
• 5703000 | Bipolar disorder in partial remission (disorder) |

Unacceptable examples,
• X in early full remission
• X in sustained full remission
• X in sustained partial remission

Conditions with associated symptoms should be expressed and modeled like combined disorders. *Due to* situations are acceptable but not simple Co-occurrent.

For example,
• 703850002 | Delirium due to benzodiazepine withdrawal (disorder) |

Concepts containing X without Y are considered on a case-by-case basis.

Acceptable example,
• 724735003 | Oppositional defiant disorder without chronic irritability-anger (disorder) |

Unacceptable example,
• Bipolar type II disorder with current episode moderately depressive without psychotic symptoms

See also relative section(s):
Remission

Multisystem disorders
Multisystem disorders are often rare conditions. There may be limited information about such disorders, so they should be carefully modeled.

When determining parent concepts:
• A multisystem parent concept should be included.
• Genetic or inherited disorders should be modeled in the same way as other genetic and inherited disorders.
• The manifestations of the disorder must always necessarily be true before assigning the relevant parents.
• Attributes must also always necessarily be true.
  • For example,
    • 702410002 | Iris coloboma with ptosis, hypertelorism, and mental retardation (disorder) |
      Since the coloboma of the iris is not always present, coloboma would not be explicitly modeled in the relationships.

Some multisystem disorders can be named by their manifestations. The FSN should be descriptive rather than just a list of names.

For example,
• 717909004 | Bilateral microtia with deafness and cleft palate syndrome (disorder) |

A multisystem disorder with an eponymous syndrome name should be included as a synonym only.

Neoplasm
Tumor vs. neoplasm
The word *tumor* has two primary meanings: a mass, regardless of whether it is neoplastic or not; or a neoplastic mass. The term *neoplasm* is preferred since it is less ambiguous than tumor. The word *tumor* is acceptable as a synonym but not as a preferred term.

For example,
Primary vs. secondary neoplastic disorders
SNOMED CT follows ICD-O and ICD-10 where secondary malignant neoplasm of site X is uniformly interpreted to mean that metastasis has occurred to site X.

For example,

- 94521000 [Secondary malignant neoplasm of rib (disorder)]

For concepts that describe metastasis from a malignant neoplasm, SNOMED CT explicitly uses the word from.

For example,

- 315006004 [Metastasis from malignant neoplasm of lung (disorder)]

In SNOMED CT, metastases are modeled with two relationship groups, each with an appropriate morphology and site.

For example,

712849003 [Primary malignant neoplasm of prostate metastatic to bone (disorder)]:
- IS A (attribute): Disease (disorder)
- Finding site (attribute): Bone structure (body structure) and Associated morphology (attribute): Neoplasm, metastatic (morphologic abnormality)
- Associated morphology (attribute): Malignant neoplasm, primary (morphologic abnormality) and Finding site (attribute): Prostatic structure (body structure)

Neoplasia
When modeling neoplasia, distinguish structure from process. Do not use neoplasia in the FSN to identify the structure (even though it implies it). Use 126537000 [Neoplasm of bone (disorder)], not neoplasia of bone.

Neoplastic disease refers to the process of neoplasia, leading to the formation of a neoplasm.

Where the definition is primary, the associated morphology: 86049000 [Malignant neoplasm, primary (morphologic abnormality)] is used.

Where the definition is primary or secondary, the morphology: 367651003 [Malignant neoplasm of primary, secondary, or uncertain origin (morphologic abnormality)] is used.

Neoplasm versus hamartoma
A neoplasm is defined as a growth of tissue no longer under normal control. A hamartoma is defined as a benign, self-limiting growth of disorganized mature cells normally found in the region, representing faulty development. SNOMED CT has disorder (and morphologic abnormality) concepts and subtypes representing neoplasia, hamartomas, and tumors.

The SNOMED CT concept 399981008 [Neoplasm and/or hamartoma (disorder)] has six subtypes:

- angiomatosis
- hamartoma
- hemangioma
- lymphangioma
- melanocytic nevus
- neoplastic disease

The SNOMED CT concept 400177003 [Neoplasm and/or hamartoma (morphologic abnormality)] also has six subtypes:

- angiomatosis
- blood vessel tumor
- hamartoma
- lymphatic vessel tumor
- melanocytic nevus
- neoplasm
Nevus

The word *nevus* has many different meanings. The differences are generally based on answers to the following questions:

- Is it necessarily on the skin? Or can it be located in mucosal sites or other sites?
- Is it necessarily visible? Or can it be in internal locations such as gastric mucosa, etc?
- Is it necessarily present at birth? Or can it occur later in life?
- Is it necessarily dark and made of melanocytes? Or can it be non-pigmented, or made of other types of cells?
- Is it necessarily made of tissue that is normally present at the site? Or can it be ectopic?
- Does it exclude benign neoplasms?

Some common meanings of nevus based on some combinations of answers to the questions are as follows:

- A birthmark, that is, any visible spot on the skin or oral mucosa present since birth, regardless of tissue of origin, excluding benign neoplasms.
- Any benign cluster of melanocytes, regardless of location, and regardless of pigmentation, whether present since birth or appearing later.
- Any cutaneous hamartoma. This excludes non-cutaneous sites, and excludes neoplasms and ectopic tissue, such as choristomas.

As a result of this wide variation in meaning, any SNOMED CT FSN containing the word *nevus* may be ambiguous. For example, the term *vascular nevus* may mean:

- Congenital blood vessel tumor in the skin
- Congenital blood vessel hamartoma or neoplasm that is visible somewhere (not only the skin, but also the mucosa, whether visible externally or not)
- Congenital blood or lymphatic vessel tumor in the skin
- Congenital blood or lymphatic vessel hamartoma or neoplasm that is visible somewhere
- Any of the above but not necessarily congenital

A better FSN for vascular nevus (morphologic abnormality) would be vascular hamartoma (morphologic abnormality). Likewise, a better FSN for congenital vascular nevus (disorder) would be congenital vascular hamartoma (disorder).

In those cases where common clinical usage of a term containing nevus is unambiguous, there is no need to inactivate the description or the concept.

Overlapping neoplasm

Overlapping neoplasm concepts refer to a neoplasm that overlaps two or more adjacent sites. For clarity, the phrase *overlapping sites* should be included in the descriptions for the FSN and PT for new overlapping neoplasm content.

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**Do not use overlapping lesion wording.**

For example,

188247000 | Malignant neoplasm of overlapping sites of bladder (disorder) |

For modeling an overlapping neoplasm concept, if the concept refers to contiguous sites involving more than one anatomical site, then a separate role group is used for each finding site.

However, assigning a role group for each finding site does not sufficiently define a concept but merely indicates the presence in both sites of a neoplasm. Where a relevant primitive existing overlapping neoplasm concept is available, this can be used as a stated primitive parent to sufficiently define the concept.

For example,

Sufficiently defined concept 721624000 | Primary adenocarcinoma of overlapping sites of esophagus (disorder) | has a stated primitive parent of 187824009 | Malignant neoplasm of overlapping sites of digestive system (disorder). |

Obstruction
Since an obstruction describes blockage inside the space of a tubular structure, the Finding site of obstruction concepts should be a value from the 113342003 [Structure of lumen of body system (body structure)] subhierarchy.

For example,

When modeling gastrointestinal tract obstruction concepts, the Finding site value should be a value from the 432899004 [Structure of lumen of gastrointestinal tract (body structure)] hierarchy as the site obstructed is the lumen of the tract.

Osteoarthritis
396275006 [Osteoarthritis (disorder)] is regarded as a degenerative disease, despite the -itis in its name. Because of this, 396275006 [Osteoarthritis (disorder)] is not a subtype of arthritis in the disorder hierarchy but instead, the more general, 399269003 [Arthropathy (disorder)]. Arthritis is inflammatory by definition, but osteoarthritis has a subclass in the medical literature called non-inflammatory osteoarthritis. In fact, according to many authoritative sources, osteoarthritis is usually regarded as a non-inflammatory disease, and therefore it is not strictly a subtype of arthritis.

Structuring the hierarchy this way does not imply that there are no cases of osteoarthritis with inflammation, nor does it rule out inflammation as an etiologic or contributory factor. It is well established that inflammation often occurs in osteoarthritis, and treatment with anti-inflammatory agents has been more effective than pure analgesics in many cases. Despite growing evidence of the role of inflammatory cytokines in osteoarthritis, it is not always necessarily an inflammatory disorder of the joint.

Pneumonia vs. Pneumonitis
The terms pneumonia and pneumonitis are often used interchangeably. In SNOMED CT, pneumonia should be used for infectious causes, and pneumonitis should be used for noninfectious causes.

Pneumonia is a type of pneumonitis, as inflammation is present in both. The distinguishing feature between the two disorders is the presence of infection in pneumonia. Pneumonia should have a pathological process of infectious process, pneumonitis should not.

Consolidation is a feature of most forms of pneumonia. It may not be a feature of some atypical pneumonias, e.g. mycoplasma pneumonia.

Except as noted above, the morphologic abnormality for 233604007 [Pneumonia (disorder)] is 707496003 [Inflammation and consolidation (morphologic abnormality)].

The morphologic abnormality for 205237003 [Pneumonitis (disorder)] is 409774005 [Inflammatory morphology (morphologic abnormality)].

Poisoning
When modeling poisoning disorders, ensure that the disorder being described is caused by the substance or active ingredient in the product selected as the causative agent (attribute) value. Do not add poisoning disorders if the causative agent is a product constituent (e.g. adjuvant, carrier, preservative, flavoring, stabilizer, or other inactive ingredient) that cannot be identified as the causative agent.

Pulmonary embolism
Pulmonary embolus (PE) refers to obstruction of the pulmonary artery or one of its branches by material (e.g. thrombus, tumor, air, or fat) that originated elsewhere in the body. When modeling embolism disorder concepts with pulmonary in the FSN, the Finding site is 782966009 [Structure of artery of pulmonary circulation (body structure)].

Remission
Disorder in remission <X> disorder in remission concepts require a stated relationship to the appropriate primitive Disorder in remission supertype, in addition to the appropriate supertype for the disorder.

For example,
16270831000119107 |Bulimia nervosa in partial remission (disorder)| has stated parents of 698698008 |Bulimia nervosa in remission (disorder)| and 765207007 |Disorder in partial remission (disorder)|.

Figure 1: Stated view of 16270831000119107 |Bulimia nervosa in partial remission (disorder)|

Where the primitive supertype for the disorder is |Disease (disorder)|, only the Disorder in remission supertype will be required.

For example, 91856007 |Acute lymphoid leukemia in remission (disorder)| has only one stated parent of 765205004 |Disorder in remission (disorder)|, because a potential supertype of 64572001 |Disease (disorder)| would be unnecessary.
Figure 2: Stated view of 91856007 |Acute lymphoid leukemia in remission (disorder)|

See also relative section(s):

Mental health

Rheumatoid arthritis
Rheumatoid arthritis (RA) is a multisystem, inflammatory, autoimmune disorder; the exact etiology is unknown. RA is a disease primarily of the joints and is clinically known as an 'arthritis' although extra-articular manifestations occur. Extra-articular features include nodules, carditis and pericarditis, vasculitis, lung disorders, and other manifestations.

69896004 |Rheumatoid arthritis (disorder)| remains a primitive concept in SNOMED CT and must be stated as a parent (ISA relationship) for all rheumatoid arthritis concepts.

For example,

201776007 |Rheumatoid arthritis of sacroiliac joint (disorder)|
Figure 1: Stated view of 201776007 |Rheumatoid arthritis of sacroiliac joint (disorder)|

Example of extra-articular rheumatoid manifestation,
28880005 |Rheumatoid arthritis with carditis (disorder)|
Figure 1: Stated view of 28880005 |Rheumatoid arthritis with carditis (disorder)|

410795001 |Juvenile rheumatoid arthritis (disorder)| has been inactivated with an inactivation reason of Outdated with a target replacement of 410502007 |Juvenile idiopathic arthritis (disorder)|. Subtypes of Juvenile idiopathic arthritis (disorder) are now modeled to reflect the up-to-date classification of this disorder.

Transplant rejection vs. failure
Modeling for transplantation rejection vs. failure is as follows:

**Transplantation rejection**
- IS A = disease
- clinical course = acute/chronic etc.
- after = transplantation of x (procedure)
- finding site = transplanted x body structure

**Transplantation failure**
- IS A = complication
- clinical course = acute/chronic etc.
- after = transplantation of x (procedure)
- finding site = transplanted x body structure

**Trauma and Injury**

Trauma, injury

There is a need to represent both traumatic and non-traumatic injuries as well as those in which it is undetermined whether the cause of the injury was due to trauma or not. There are forms of trauma that do not result in structural damage, such as emotional trauma, which are defined as traumatic injuries. Concepts that do not specify trauma are now modeled with <<Damage (morphologic abnormality) but are not necessarily assigned a Due to (attribute) of Traumatic event (event), unless the form of damage can only incur morphologic trauma. In other words, if a concept refers to an injury, and that injury may occur either through trauma or non-traumatic means (e.g. tumor, ischemia, etc.), then it should be modeled without a Due to (attribute) of [Traumatic event]. If, however, the term does not specify trauma, but the type of injury can only occur as a result of trauma (e.g. open wounds), then these concepts would have the DUE TO attribute added.

19130008 |Traumatic abnormality (morphologic abnormality)| has been inactivated effective January 2021 in order to separate mechanism of injury (i.e. trauma) from structure (i.e. damage). Historically, injury concepts have been modeled in SNOMED CT as damage to a body structure, unless specifically stated as non-traumatic.

- Traumatic injuries are now being modeled as damage to a body structure due to traumatic event.
- Non-traumatic injuries are being remodeled as damage to a body structure but without a Due to (attribute) relationship to Traumatic event (event). Nontraumatic injuries should be modeled as a subtype of 1119219007 |Nontraumatic injury (disorder)|.

417163006 |Traumatic or non-traumatic injury (disorder)| is currently modeled with GCIs to reflect the two notions of damage without trauma (non-traumatic injury) and trauma with or without damage (traumatic injury).

Friction injury, abrasion

An injury due to friction can be represented using 400152004 |Friction injury (morphologic abnormality)|, in which case it will not classify as a kind of wound.

For example,
- 47222000 |Friction injury of tooth (disorder)|
- 400068007 |Mechanical irritation (morphologic abnormality)|
However, most disorders that are named *abrasion* imply that skin or other tissue has been abraded (scraped or worn away). Thus, they are also considered wounds and will correctly classify as wounds after assigning the correct morphology, 400061001 |Abrasion (morphologic abnormality)|.

For example,

- 211039006 |Abrasion, chest wall (disorder)|

**Rupture**

Ruptures are modeled with an |Associated morphology (attribute)| of 125671007 |Rupture (morphologic abnormality)|. A disorder concept modeled with a Rupture (morphologic abnormality) will classify as a subtype of 417163006 |Traumatic or non-traumatic injury (disorder)|.

- Traumatic rupture concepts are modeled with a |Due to (attribute)| of << |Traumatic event (event)|
- Nontraumatic rupture concepts are modeled as a subtype of 1119219007 |Nontraumatic injury (disorder)|

**Effective July 2021**

Both 415747007 |Traumatic rupture (morphologic abnormality)| and 125672000 |Nontraumatic rupture (morphologic abnormality)| are inactive effective from July 2021.

**Vaccine-related overdose**

For the January 2020 Release, vaccine-related overdose concepts in the Clinical Finding/Disorder hierarchy were inactivated. They were replaced with concepts in the Event hierarchy, see 788094008 |Excessive dose of vaccine administered (event)| and subtypes.

When authoring, determine whether the concept describes an overdose, which is a *disorder*, or the administration or ingestion of an excessive dose, which is an *event*.

**Disorder Combination Modeling**

Many disorders can occur in combination within the same patient. Guidance on the modeling and terming of FSNs for disorder combinations aims to achieve consistency. Clinically significant disorder combinations are represented in SNOMED CT by a single concept so that users can document temporal (timing) and causal (cause/effect) relationships between the conditions.

Not all disorders occurring in combination should be precoordinated into a single concept. Conditions should be recorded as separate concepts if one can be resolved separately from the other, or if the two conditions require separate management or treatment plans. Multiple clinical conditions should not be precoordinated in order to facilitate convenient recording in the medical record, even if the two conditions are often reported together.

For example,

- The clinical conditions *gastroenteritis* and *dehydration* often occur in combination but require separate resolution, and therefore, are best recorded separately in the medical record as 25374005 |Gastroenteritis (disorder)| and 34095006 |Dehydration (disorder)|.

To express an association between conditions, one of the following associations is used:

- Simple co-occurrence: two or more conditions have no direct causal or temporal relationship but are found together more often than by random chance
- Causation 1: the cause is another finding or disorder, an event, or procedure
- Causation 2: the cause is a physical force, physical object, organism, or substance
- Temporal association: the timing of the two conditions occur before, during, or after each other

**Causality**

Causal relationships between disorders are represented using the Due to (attribute).

- When modeling a *disorder due to another disorder*, do not use |Complication (disorder)| as a supertype.
When modeling a *disorder due to a surgical procedure*, use |Complication (disorder)| as a supertype.

**Truth Table**

When considering disorder combinations two questions can be asked:

1. Is there a *causal* relationship?
2. What is the *temporal* relationship?

The following truth table provides the possible combinations/answers. It allows authors to assign combination disorders to one category, to which the appropriate modeling and FSN construction is applied. The stricter rules for FSN construction do not prevent the addition of more familiar connectives in other descriptions, for example *with, or associated with*.

<table>
<thead>
<tr>
<th>Is there a stated causal relationship?</th>
<th>X due to Y</th>
<th>X caused by Y</th>
<th>X and Y should be documented separately</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yes, the cause is another finding or disorder, an event, or procedure. This is causation 1.</td>
<td>X due to Y</td>
<td>X caused by Y</td>
<td>X and Y should be documented separately</td>
</tr>
<tr>
<td>Yes, the cause is a physical object or force, organism, or substance. This is causation 2.</td>
<td>X due to Y</td>
<td>X caused by Y</td>
<td>X and Y should be documented separately</td>
</tr>
<tr>
<td>No</td>
<td>X follows Y</td>
<td>X after Y</td>
<td>N/A</td>
</tr>
<tr>
<td>X precedes Y</td>
<td>X due to Y</td>
<td>X before Y</td>
<td>N/A</td>
</tr>
<tr>
<td>X occurs during Y</td>
<td>X due to Y</td>
<td>X during Y</td>
<td>N/A</td>
</tr>
<tr>
<td>X occurs before, during, and/or after Y</td>
<td>X due to Y</td>
<td>X temporally related to Y</td>
<td>N/A</td>
</tr>
</tbody>
</table>

Note: *Temporally related to (attribute)* and its subtypes *Before* and *During* are only approved to model perioperative complications.

<table>
<thead>
<tr>
<th>Simple Co-occurrence</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Modeling pattern</strong></td>
</tr>
</tbody>
</table>
Assign each condition as a supertype (or ensure that each participating disorder is present in the ancestor tree following classification).

Use simple co-occurrence for two or more conditions that are strongly associated by means other than causality or a temporal relationship (e.g. a common predisposition) where representing such conditions as separate statements would result in a loss of the associated between the conditions.

For example, named syndromes, such as 398114001 | Ehlers-Danlos syndrome (disorder) |
manifestations of systemic disorders, such as 83901003 | Sjögren's syndrome (disorder) |

Do not use simple co-occurrence for those disorders with more than one anatomical site or more than one associated morphology. Those disorders should rather be represented as individual concepts in a medical record.

FSN: X with Y

PT: Eponyms may be used if acceptable.

Examples:
Sinusitis with nasal polyps (disorder)
Acute bronchitis with bronchiectasis (disorder)

Incorrectly named legacy examples (not to be repeated):
Psoriasis-eczema overlap condition (disorder)
Hay fever with asthma (disorder)

Causation 1

There are complications which likely exist prior to a disorder or procedure.
For example, the legacy term, 609454008 | Induced termination of pregnancy complicated by acute necrosis of liver (disorder) |

- Acute necrosis of liver is the complication however, temporally it is neither due to nor during. It was likely to be present prior to the procedure. The concept will be inactivated. Instead the separate concepts 714812005 | Induced termination of pregnancy (procedure) | and 197269008 | Acute necrosis of liver (disorder) | should be documented in the medical record.

⚠️ Pattern Variance

- The FSN submitted by a requestor may be used as preferred term even if it does not comply with the above recommended pattern. However, do not use phrases such as secondary to, as a result of, etc. in lieu of due to.
- Exceptions may exist to the above policy which will need to be reviewed on a case-by-case basis.

<table>
<thead>
<tr>
<th>Causation 1</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cause is another finding, disorder, event, or procedure</td>
</tr>
<tr>
<td>Modeling pattern</td>
</tr>
</tbody>
</table>
Causation 1

For a condition caused by a clinical finding/disorder

- Ensure two conditions are represented in the supertypes and/or axioms
- Assign the causal disorder as the target of a *Due to* relationship

For a condition caused by a procedure

- For surgical procedures:
  - Ensure the caused condition is represented as a supertype and/or axiom
  - Ensure Complication (disorder) is a supertype
  - Assign the procedure as the target of a *Due to* relationship
- For non-surgical procedures:
  - Ensure the caused condition is represented as a supertype and/or axiom
  - Do not use Complication (disorder) as a supertype
  - Assign the procedure as the target of a *Due to* relationship

For a condition caused by an event

- Ensure the caused condition is represented as a supertype and/or axiom
- Assign the event as the target of a *Due to* relationship

Examples:

<table>
<thead>
<tr>
<th>Concept ID</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>735200002</td>
<td>Absence of lower limb due to diabetes mellitus (disorder)</td>
</tr>
<tr>
<td>735621005</td>
<td>Adhesions due to endometriosis (disorder)</td>
</tr>
</tbody>
</table>

Incorrectly named legacy examples (not to be repeated):

- Neutropenia associated with acquired immunodeficiency syndrome (disorder)
- Dilated cardiomyopathy secondary to granuloma (disorder)

There are approximately 430 legacy concepts with ‘co-occurrent and due to’ in the description. Do not add new concepts with the terming ‘co-occurrent and due to’, instead use co-occurrence modeling (both conditions are represented in a supertype) in addition to the *Due to* (attribute) if warranted by the clinical condition.

☑ **Umbilical cord complication**

- Model as IS A 362972006 *Disorder of labor / delivery (disorder)* due to X (disorder). The concept 48287005 *Umbilical cord complication (disorder)* is to be inactivated.
### Causation 2

Causation 2 is when 1 the cause is a material entity, and 2 the means of exposure/introduction are not significant.

1. A material entity refers to a concept within the Substance, Physical object, Pharmaceutical/biologic product, Physical force, and Organism hierarchies.
2. If the means of exposure/introduction are significant, then the causal factor is represented by a concept from the Event hierarchy, and the concept is modeled as Causation 1.

<table>
<thead>
<tr>
<th>Modeling pattern</th>
<th>Naming pattern</th>
</tr>
</thead>
<tbody>
<tr>
<td>Assign the caused disorder (X) as a supertype, or ensure that the caused disorder is a supertype following classification</td>
<td>X caused by Y</td>
</tr>
<tr>
<td>Assign the causal factor (Y) as the value of a Causative agent (attribute)</td>
<td></td>
</tr>
</tbody>
</table>

**Example:**

<table>
<thead>
<tr>
<th>Code</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>291000119100</td>
<td>Contact dermatitis caused by chemical (disorder)</td>
</tr>
</tbody>
</table>

Incorrectly named legacy examples (not to be repeated):

- Choking due to airways obstruction (finding)
- Coma associated with diabetes mellitus (disorder)
- Laser-induced burn (disorder)

### Temporal sequencing without causation

Modeling captures and emphasizes *non-overlap*. Explicit causation may be captured using both *due to* AND *after* relationships, as in the truth table above.

<table>
<thead>
<tr>
<th>Modeling pattern</th>
<th>Naming pattern</th>
</tr>
</thead>
</table>
## Temporal sequencing without causation

Assign the condition or procedure that occurred first in the patient as the target of an After (attribute) relationship. Assign the condition that occurred second as a supertype (or ensure its presence in the ancestor tree).

- **For example,**
  
  402490007 |Calcinosis following localized fat necrosis (disorder)|

  The fat necrosis occurred first in the patient, so this concept will have an After (attribute) with a value of Fat necrosis (disorder). The calcinosis occurred secondarily, and thus Calcinosis (disorder) is a supertype of this concept.

For disorders with a temporal relationship to a surgical procedure, assign an additional supertype of 116223007 |Complication (disorder)|.

- **For example,**

  16055031000119100 |Astigmatism of right eye following operative procedure (disorder)|

  The operative procedure occurred first in the patient, so this concept will have an After (attribute) with a value of Surgical procedure (procedure). The astigmatism occurred secondarily, so Astigmatism (disorder) is a supertype of this concept. Because this disorder has a temporal relationship to a surgical procedure, an additional supertype of Complication (disorder) is assigned.

### Where X occurs after Y:

- if it is not specified that X is due to Y (although causality is frequently implied), construct the FSN as X following Y
- if it is specified that X is due to Y, construct the FSN as X due to and following Y

## Associated with (attribute)

In general, 47429007 |Associated with (attribute)| should be avoided due to the ambiguity which it conveys and the difficulty in applying this role consistently. Instead, Due to is used when there is a direct causal relationship between the conditions; otherwise, the clinical conditions should be recorded as separate concepts in the medical record.

There are a couple of exceptions when the use of 47429007 |Associated with (attribute)| is appropriate:

1. **General grouping concepts which aggregate more specific associations**
   
   e.g. 6211002 |Polyarthritis associated with another disorder (disorder)| subsumes two children
   
   201972000 |Allergic arthritis of multiple sites (disorder)| modeled with 42752001 |Due to (attribute)| of 419076005 |Allergic reaction (disorder)|
   
   422565003 |Post-infective polyarthritis (disorder)| modeled with 255234002 |After (attribute)| of 40733004 |Infectious disease (disorder)|

2. **Device infections, i.e an infection of the tissue surrounding an implanted or inserted device, not due to the device itself.**
   
   - Associated with is used to associate the device with the infection.

3. **Intolerance to substances, i.e the propensity of an adverse reaction to a substance to occur (other than hyper-sensitivity or allergic or non-allergic hypersensitivity).**

4. **There is no intolerance process that serves as the value for Has realization.**
   
   - Associated with is used to associate the intolerance to the substance.

### Is cause a disorder or material entity?

It must be determined if a disorder is caused either by another disorder or by a material entity. A material entity is a concept found in Substance, Physical object, Pharmaceutical/biologic product, Physical force, or Organism subhierarchies. These subhierarchies are the current range constraints for the Causative agent (attribute) in the...
Clinical finding domain. For combined disorders where a cause can be either a disorder (eg, alcoholism) or a material entity (eg, alcohol):

- Model as *due to disorder* if it is the indirect cause.
  
  For example,
  
  - Megaloblastic anemia *due to alcoholism* (disorder)

- Model as *caused by material entity* if it is the direct cause.
  
  For example,
  
  - Inflammation of pancreas *caused by alcohol* (disorder)

Is cause a disorder or infectious organism?

In modeling concepts related to infectious diseases, a number of considerations need to be taken into account.

Firstly, when the disorder is an infectious disease itself, and the organism is specified, then the concept will be modeled with

- |Causative agent (attribute)| with the specified organism
- |Pathological process (attribute)| with the type of infectious process

Secondly, Disorders can be modeled with |Due to|, |After| or |Due to| and |After| relationships to infectious diseases.

Note If the focus disorder is itself an infectious disorder, it will also have a |Causative agent| relationship when the organism is specified.

Examples

- |Causative agent| relationship: 721742004 |Otitis media caused by Streptococcus pneumoniae (disorder)|
- |Due to| relationship: 698733009 |Intestinal obstruction due to tuberculosis (disorder)|
- |Due to| and |Causative agent| relationship: 866044006 |Mycosis due to human immunodeficiency virus infection (disorder)|
- |After| relationship: 182961000119101 |Acute disseminated encephalomyelitis following infectious disease (disorder)|
- |After| and |Causative agent| relationship: 4740000 |Herpes zoster (disorder)|
- |Due to| and |After| relationship: 1148594002 |Chronic arthritis due to and following rheumatic fever (disorder)|
- |Due to| and |After| and |Causative agent| relationship: 15992311000119100 |Keratitis of left eye due to herpes zoster (disorder)|

⚠️ *Applying the |Due to|, |After| or |Due to| and |After| relationships to a concept will not lead to it being a subtype of |Infectious disease (disorder)| unless it is itself an infectious disease.*

Exception to naming convention for combined disorders

Exceptions may exist to the above guidance which requires review on a case-by-case basis.

The FSN submitted by a requestor may be used as preferred term even if it does not comply with the above recommended pattern. However, do not use phrases such as *secondary to, as a result of*, etc. in lieu of *due to*.

Rather than the naming conventions described above, use the names that are accepted clinical parlance and that represent specific pathophysiologic entities for some combined disorders, as the preferred term.
### Disorder combination modeling

- Covers combinations of only two disorders. However, combinations often include more than two disorders (for example, syndromes). Document multiple conditions in a single statement only for syndromes or strong associations based on a common predisposing factor.
- Does not cover absent components or negation
- Does not cover cases where combination concepts are demonstrably classification-derived (This limitation accepts that some content may be so obviously based on a class or category in a classification that it would be undesirable to reinterpret its semantics.)
- The modeling approach may be difficult to apply in all cases of combined disorders; domain-specific templates should be developed to ensure modeling consistency and accuracy.

(See also, Appendix, Concept Models: Disorder Combinations)

Heuristics for Co-occurrent Genomic Disorders

Germline chromosomal abnormality co-occurring and causing disorder: 41040004 | Complete trisomy 21 syndrome (disorder)
If the phenotype is always caused by a specific genotype, there is no need to include the cause in the FSN or clarify with a Due to relationship.

Germline nucleotide sequence variant co-occurring and causing disorder: 190905008 | Cystic fibrosis (disorder)
Modeling for germline mutations causing conditions, such as cystic fibrosis, should have mutations, Occurrence = congenital, and Due to (attribute) the mutation finding.

For example,
- Cystic fibrosis due to G542X mutation

Somatic NSV (NCBI structural variant) co-occurring and poly-etiologic: BRAF V600E positive melanoma
Somatic mutations leading to cancer, such as malignant melanoma with BRAF V600E mutation, should have dual supertypes, including the malignant disorder and the somatic mutation, and Due to (attribute) with the associated somatic mutation finding.

For example,
- Melanoma with BRAF V600E mutation

Somatic IHC (immunohistochemical) finding co-occurring but not etiologic: Estrogen-receptor status in breast cancer
Representing two associated findings in a single concept may be convenient for recording; however, the representation of the two notions should be recorded separately.

For example,
- Breast cancer occurring with positive estrogen-receptor assay should be recorded in the information model as two separate concepts

The term phrase, "co-occurrent and due to" is no longer to be used in the fully specified name. There are existing concepts that use the co-occurent and due to pattern, but these will be re-termed. Genetic mutations that cause a disorder are by definition co-occurent, so there is no need to represent this in the FSN, but they should be modeled as co-occuring, i.e. supertypes for both conditions should be present.
Complication and Sequela Modeling

Combined disorders can occur, for example:

- One disorder causes the other (causal relationship)
- One disorder is temporally related to another
- Two disorders have both a causal and temporal relationship to each other

Attributes that can be used to define such causal and temporal relationships are:

- Associated with (attribute)
- Causative agent (attribute)
- Due to (attribute)
- Temporally related to (attribute)
  - Before (attribute)
  - During (attribute)
  - After (attribute)

Causal and temporal attributes are used in modeling complications and sequelae. Due to the various combinations of cause and timing, disorders may occur as both a complication and as a sequela. The words complication and sequela may or may not be in the FSN description.

Complication

A **complication** is a disorder due to or temporally related to <<Surgical procedure (procedure) or an event that is neither a natural progression nor an expected outcome of its cause. A complication parent is only assigned for surgical complications. Surgical complications may be:

due to surgery,

be temporally related to surgery or

be due to and temporally related to surgery.

The phrase "Complication of" implies causality, and thus “due to” should be used in the description.

116223007 |Complication (disorder)| is no longer used as a supertype for a disorder caused by another disorder, nor for disorders following non-surgical/non-operative medical procedures.

Perioperative complications

Perioperative complications refer to complications temporally related to a surgical procedure. They include pre-operative, intra-operative and post-operative complications and are modeled with a parent of Complication (disorder) and a relationship consisting of Temporally related to (attribute) or an appropriate subtype with a value of 387713003 |Surgical procedure (procedure)|. A temporal complication does not necessarily imply a causal (Due to) relationship to the surgery itself, as the complication may be related to any disorder, event, or procedure occurring either prior, during and/or after surgery. For this reason, perioperative complications do not have a stated Due to relationship unless an underlying cause is clearly stated in the FSN.

The following naming convention applies to those conditions that occur temporally, i.e. either before, during, or after the operative episode, but do not have a causal relationship.

**FSN:** Postoperative X (disorder)

**PT:** Postoperative X

For example,

- Perioperative hematoma (disorder)
- Postoperative hypothyroidism (disorder)

The following attributes are used in the various combinations of complication and sequela modeling:

**After**

**After without causal relationship**
This attribute is used to model concepts in which a clinical finding occurs after another clinical finding, procedure, or event. Neither asserting nor excluding a causal relationship, it instead emphasizes a sequence of events. Naming pattern is ‘x following y’.

For example,

123948009 |Disorder following viral disease (disorder)| occurs After 34014006 |Viral disease (disorder)|

Figure 1: Stated view of 123948009 |Disorder following viral disease (disorder)|

Post-infectious disorders are not subtypes of infectious disorders. The After attribute is used for linking post-infectious disorders with their associated infections.

After with causal relationship
The Due to and After attributes are used to model a disorder that occurs after a disorder or procedure with a causal relationship. Both the cause and the After relationship must be specified. The naming pattern is ‘due to and following’.

Before
This attribute is used to model a preoperative complication. Strictly, a preoperative complication is a disorder that complicates the procedure, rather than being a complication of that procedure. A preoperative complication might be considered to be a disorder that exists prior to surgery that adversely affects the surgery or that results in an intraoperative or postoperative complication.

During
During without causal relationship
This attribute is used to model a disorder that occurs during a procedure. Only use this attribute when explicit in the FSN (e.g. *Intraoperative complication* or *X complication during Y surgical procedure*).

For example,

![Diagram](image)

Figure 2: Stated view of 713890008 |Hypoxemia during surgery (disorder)|

During with causal relationship
Due to and During attributes can be used to model a disorder that occurs during a procedure (e.g. intraoperative complication), with a causal relationship. Both a cause and a temporal relationship to the cause must be specified.

Sequelae
A *sequela* is a disorder that is a consequence, but not an unexpected outcome, that follows after another disorder, procedure, or event. These conditions are often described with the words *following, after, post, or sequela(e)*. ICD uses the phrase *late effects.*
Sequela can be in the following forms:

- Following
- Due to and following/after
- During and following/after

These conditions should be modeled with After (and also Due to if there is a causal relationship).

For example,

Disorder due to and following another disorder = 698737005 |Obstructive hydrocephalus due to and following meningitis (disorder)|

Figure 3: Stated view of 698737005 |Obstructive hydrocephalus due to and following meningitis (disorder)|
Naming conventions for sequelae
FSN: Disorder X [due to and] following <<disorder /<<procedure /<event
PT: Disorder X [due to and] following <<disorder /<<procedure /<event
SYN: [Disorder X as a] Sequela of <<disorder /<<procedure /<event
SYN: [Disorder X as a] Late effect of <<disorder /<<procedure /<event

For example,
• Disorder due to and following another disorder (disorder)
• Disorder due to and following meningitis (disorder)
• Disorder due to and following procedure (disorder)

Naming conventions for surgical sequelae (temporal relationship but no causal relationship)
Not all surgical sequelae are complications of surgery but rather expected late effects.
FSN: Disorder X following <<387713003 |Surgical procedure (procedure)
PT: Disorder X following <<387713003 |Surgical procedure (procedure)

For example,
• Contraction of eye socket following enucleation (disorder)
• Scar following surgery (disorder)

Naming conventions for surgical sequelae complications (temporal relationship and causal relationship)
Complications that occur following surgery, but not necessarily Due to the surgery, are modeled only with an After relationship.
Complications that occur following surgery and are explicitly stated as causal/due to are modeled with a parent of 116223007 |Complication (disorder)| and with |after| |surgical procedure|.
FSN: Disorder X due to and following <<387713003 |Surgical procedure (procedure)
PT: Disorder X due to and following <<387713003 |Surgical procedure (procedure)|

For example,
• Encephalopathy due to and following cardiopulmonary bypass (disorder)
• Cataract lens fragments in vitreous of eye due to and following cataract surgery (disorder)
• Disorder due to and following breast reduction (disorder)

Environment and Geographical Location

<table>
<thead>
<tr>
<th>Environment: types of environments</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Environment: types of environments</td>
<td>• 398156002</td>
</tr>
<tr>
<td>• Location: named locations such as countries, states, or regions</td>
<td>• 223565009</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Event</th>
</tr>
</thead>
<tbody>
<tr>
<td>Occurrences impacting health or health care; not procedures or interventions</td>
</tr>
<tr>
<td>Examples</td>
</tr>
<tr>
<td>• 242039002</td>
</tr>
<tr>
<td>• 405621004</td>
</tr>
</tbody>
</table>
Event Attributes Summary

When authoring in this domain, these are the approved attributes and allowable ranges. They are from the Human Readable Concept Model (HRCM).

Domain Information for 272379006 | Event (event) |
--- | --- |
Domain Constraint | << 272379006 | Event (event) |
Parent Domain | - |
Proximal Primitive Constraint | << 272379006 | Event (event) |
Proximal Primitive Refinement | - |

Author View of Attributes and Ranges for 272379006 | Event (event) |
| Attribute | Grouped Cardinality | In Grouped Cardinality | Range Constraint |
| 255234002 | After (attribute) | 1 | 0..* | 0..1 | << 272379006 | Event (event) | OR | << 404684003 | Clinical finding (finding) | OR | << 71388002 | Procedure (procedure) |
| 47429007 | Associated with (attribute) | 1 | 0..* | 0..* | << 105590001 | Substance (substance) | OR | << 260787004 | Physical object (physical object) | OR | << 272379006 | Event (event) | OR | << 404684003 | Clinical finding (finding) | OR | << 410607006 | Organism (organism) | OR | << 71388002 | Procedure (procedure) | OR | << 78621006 | Physical force (physical force) |
| 288556008 | Before (attribute) | 1 | 0..* | 0..1 | << 71388002 | Procedure (procedure) |
| 246075003 | Causative agent (attribute) | 1 | 0..* | 0..1 | << 105590001 | Substance (substance) | OR | << 260787004 | Physical object (physical object) | OR | << 373873005 | Pharmaceutical / biologic product (product) | OR | << 410607006 | Organism (organism) | OR | << 78621006 | Physical force (physical force) |
| 42752001 | Due to (attribute) | 1 | 0..* | 0..1 | << 272379006 | Event (event) | OR | << 404684003 | Clinical finding (finding) | OR | << 71388002 | Procedure (procedure) |
| 371881003 | During (attribute) | 1 | 0..* | 0..1 | << 71388002 | Procedure (procedure) |
| 246454002 | Occurrence (attribute) | 1 | 0..* | 0..1 | << 282032007 | Periods of life (qualifier value) |
| 726633004 | Temporally related to (attribute) | 1 | 0..* | 0..* | << 404684003 | Clinical finding (finding) | OR | << 71388002 | Procedure (procedure) |

⚠️ Authoring guidelines for the use of attributes in the Event hierarchy are being established.
Event Modeling

When modeling an event, be sure to model the event itself and not the outcome of an event. The outcome of an event would be a finding or a disorder.

Event attributes

Under review at this time.

The Event hierarchy should not precoordinate periods of life/occurrence within the event concept.

---

Allowed ranges

The allowed ranges will not prevent some incorrect modeling. Some allowed attributes have not yet been used for modeling in the Event domain. The planned QI project will review the modeling to ensure consistency of use the allowed attributes.

---

Modeling

**Vaccine-related overdose**

- For the January 2020 Release, vaccine-related overdose concepts in the Clinical Finding/Disorder hierarchy were inactivated. They were replaced with excessive dose concepts in the Event hierarchy.
- When authoring, determine whether the concept describes an overdose, a disorder, or the administration or ingestion of an excessive dose, an event.

---

Observable Entity

<table>
<thead>
<tr>
<th>Definition</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Information about a quality/property to be observed and how it will be observed</td>
<td>• 416540001 Calcium deposit observable (observable entity)</td>
</tr>
<tr>
<td></td>
<td>• 276885007 Core body temperature (observable entity)</td>
</tr>
</tbody>
</table>

Observable Entity vs. Evaluation Procedure

The observable entity and evaluation procedure hierarchies have some of the same attributes. There is not and should not be a one-to-one correspondence between the two hierarchies.

At this time, SNOMED CT contains some concepts in the evaluation procedure hierarchy which logically belong in the observable entity hierarchy. This is a legacy problem that continues to cause confusion. These concepts will move to the observable entity hierarchy as part of the QI project in the future. In addition, if we identify existing duplicate concepts between the two hierarchies, this will also be corrected. Concepts will not be duplicated between the observable entity hierarchy and procedure hierarchy, and requests for such will not be added.

In response to requests for an observable entity concept when the procedure concept exists, create an observable entity concept and inactivate the procedure concept.

While some users have indicated they want to use a procedure concept for ordering a test and an observable concept for reporting the result, this is not an acceptable use case. An evaluation procedure being ordered implies that there is an expectation that a value, in association with the ordered procedure will be provided. Evaluation procedures, for all intents and purposes, are observables with another semantic tag. The nature of their top level parent (Evaluation procedure) implies that they require a value in order to be assessed. Thus they can be used equivalently with observables.

As for the progression of the completion of an assessment, that is related to the state diagram (i.e., status) of the progression of a procedure and should not be precoordinated, but handled by the information system in which
orders are processed (it is dynamic, not static). The information system should be able to capture the status of a procedure (e.g., ordered, in process, completed). We would not expect the terminology to pre-coordinate this.

As an example, LOINC recognizes that there are three different aspects to an observable: 1) those that can serve as both an order and an observation (e.g. blood glucose level); 2) those that can be ordered but not directly resulted (e.g. urinalysis, which is a convenience order for multiple individual observations on urine); and 3) those that can only be resulted and not directly ordered (usually part of an automated system, such as computation of MCHC in hematology). LOINC assigns this aspect with an attribute value. It is not one of the six main LOINC parts typically visible to users, however it is included in the LOINC database.

### Use of Observable Entities

Observables entities may be used to:

- Code elements on a checklist or assign values to elements.
  - For example, *color of nail* is an observable entity. *Gray nails* is a finding.

- Code headers on a template
  - For example, the observable entity, gender, may be used to code a section of a template titled gender. The user would choose masculine, feminine, transgender, etc. which would then constitute a finding such as 703117000 |Masculine gender (finding)|.

### Types of Observable Entities

There are four general types of observable entities for use in health care. Each has different representation requirements and patterns, i.e. the set of attributes will vary.

- **Quality.** A characteristic, feature, or property that is inherent in someone or something.
  - For example, mass of a person, temperature of internal organs, concentration of sodium in plasma, angle of a joint

- **Disposition.** A characteristic or feature that is not always realized in full.
  - For example, antibiotic susceptibility of a certain population

- **Function.** The ability of a person, some part of a person, or a thing to perform activities or realize processes.
  - For example, ability to walk

- **Process.** A process or outcome of a process
  - For example, secretion rate, heart rate, respiratory rate

Some areas of the observable entity hierarchy need clarification and remodeling. This includes upper level concepts as well as leaf nodes. Notably, the content currently included in the 246464006 |Function (observable entity) subhierarchy needs to be clarified and potentially remodeled. In addition, the content currently included in the 415178003 |Process (observable entity)| subhierarchy needs review for inactivation and replacement in the 719982003 |Process (qualifier value)| hierarchy so these processes can be used as values of attributes to define observable entity concepts, e.g., via 704321009 |Characterizes (attribute)|.

### Observable Entity Attributes Summary

When authoring in this domain, these are the approved attributes and allowable ranges. They are from the Human Readable Concept Model (HRCM).
### Domain Information for 363787002 (Observable entity (observable entity))

<table>
<thead>
<tr>
<th>Domain Constraint</th>
<th>&lt;&lt; 363787002 (Observable entity (observable entity))</th>
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</thead>
<tbody>
<tr>
<td>Parent Domain</td>
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<tr>
<td>Proximal Primitive Constraint</td>
<td>&lt;&lt; 363787002 (Observable entity (observable entity))</td>
</tr>
<tr>
<td>Proximal Primitive Refinement</td>
<td>-</td>
</tr>
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</table>

### Author View of Attributes and Ranges for 363787002 (Observable entity (observable entity))

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Grouped</th>
<th>Cardinality Grouped</th>
<th>Cardinality</th>
<th>Range Constraint</th>
</tr>
</thead>
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<td>704321009 Characterizes (attribute)</td>
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<td>0..1</td>
<td>&lt;&lt; 71388002 Procedure (procedure) OR &lt;&lt; 719982003 Process (qualifier value)</td>
</tr>
<tr>
<td>246093002 Component (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
<td>&lt;&lt; 105590001 Substance (substance) OR &lt;&lt; 123037004 Body structure (body structure) OR &lt;&lt; 123038009 Specimen (specimen) OR &lt;&lt; 260787004 Physical object (physical object) OR &lt;&lt; 373873005 Pharmaceutical / biologic product (product) OR &lt;&lt; 410607006 Organism (organism) OR &lt;&lt; 419891008 Record artifact (record artifact)</td>
</tr>
<tr>
<td>704327008 Direct site (attribute)</td>
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<td>0..*</td>
<td>0..1</td>
<td>&lt;&lt; 105590001 Substance (substance) OR &lt;&lt; 123037004 Body structure (body structure) OR &lt;&lt; 123038009 Specimen (specimen) OR &lt;&lt; 260787004 Physical object (physical object) OR &lt;&lt; 373873005 Pharmaceutical / biologic product (product) OR &lt;&lt; 410607006 Organism (organism) OR &lt;&lt; 419891008 Record artifact (record artifact)</td>
</tr>
<tr>
<td>71972006 Has realization (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
<td>&lt;&lt; 719982003 Process (qualifier value)</td>
</tr>
<tr>
<td>718497002 Inherent location (attribute)</td>
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<td>0..*</td>
<td>0..1</td>
<td>&lt;&lt; 105590001 Substance (substance) OR &lt;&lt; 123037004 Body structure (body structure) OR &lt;&lt; 123038009 Specimen (specimen) OR &lt;&lt; 260787004 Physical object (physical object) OR &lt;&lt; 373873005 Pharmaceutical / biologic product (product) OR &lt;&lt; 410607006 Organism (organism) OR &lt;&lt; 419891008 Record artifact (record artifact)</td>
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<tr>
<td>704319004 Inheres in (attribute)</td>
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<td>0..1</td>
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<td>405815000 Procedure device (attribute)</td>
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<td>SNOMED CT Code</td>
<td>Description</td>
<td>Cardinality</td>
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<td>-------------</td>
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<td>Process extends to (attribute)</td>
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<td></td>
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<td>&lt;&lt; 719982003 Process (qualifier value) OR</td>
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</tr>
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<td>Property (attribute)</td>
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<tr>
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<td>Relative to (attribute)</td>
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<td>&lt;&lt; 123038009 Specimen (specimen) OR</td>
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<td>&lt;&lt; 260787004 Physical object (physical object) OR</td>
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<td>Relative to part of (attribute)</td>
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<td>&lt;&lt; 123037004 Body structure (body structure) OR</td>
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<td>&lt;&lt; 373873005 Pharmaceutical / biologic product (product) OR</td>
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<td></td>
<td>&lt;&lt; 419891008 Record artifact (record artifact)</td>
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<tr>
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<td>1 0..1 0..1</td>
<td>&lt;&lt; 117362005 Nominal value (qualifier value) OR</td>
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<td></td>
<td></td>
<td></td>
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<td></td>
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<td></td>
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<td></td>
<td>&lt;&lt; 117364006 Narrative value (qualifier value) OR</td>
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<td></td>
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<td></td>
<td>&lt;&lt; 117365007 Ordinal OR quantitative value (qualifier value) OR</td>
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<tr>
<td>246501002</td>
<td>Technique (attribute)</td>
<td>1 0..* 0..1</td>
<td>&lt;&lt; 254291000 Staging and scales (staging scale) OR</td>
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<td>&lt;&lt; 272394005 Technique (qualifier value) OR</td>
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</tr>
<tr>
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<td>Time aspect (attribute)</td>
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<td>&lt;&lt; 7389001 Time frame (qualifier value) OR</td>
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</tr>
<tr>
<td>704320005</td>
<td>Towards (attribute)</td>
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<td>&lt;&lt; 105590001 Substance (substance) OR</td>
<td></td>
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<td>&lt;&lt; 123037004 Body structure (body structure) OR</td>
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<tr>
<td>246514001</td>
<td>Units (attribute)</td>
<td>1 0..1 0..1</td>
<td>&lt;&lt; 767524001 Unit of measure (qualifier value)</td>
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<tr>
<td>424226004</td>
<td>Using device (attribute)</td>
<td>1 0..* 0..1</td>
<td>&lt;&lt; 49062001 Device (physical object) OR</td>
<td></td>
</tr>
</tbody>
</table>

**Observable Entity Defining Attributes**

**Characterizes**

This attribute specifies the process which the property describes, and on which the property (of this observable) depends. The process can be very general (e.g. excretion).
For example,

- Mass concentration ratio of silver to creatinine in 24-hour urine (observable entity) has 704321009 | Characterizes (attribute) | of excretion process
- 789098002 | Estimated quantity of intake of potassium in 24 hours (observable entity) | has
  a 704321009 | Characterizes (attribute) | of administration of substance

**Component**
This attribute is used to specify the numerator of a relational property types, e.g. ratio, concentration.

For example,

- Arbitrary concentration of Varicella-Zoster virus (observable entity) has the 246093002 | Component (attribute) | of Human herpesvirus 3

**Direct site**
This attribute is used to specify the entity on which the observation is directly made. It may also be used when the observation is indirect, i.e. when a direct observation cannot be done.

For example,

- 415974002 | Core body temperature measured at tympanic membrane (observable entity) | has the
  704327008 | Direct site (attribute) | of 42859004 | Tympanic membrane structure (body structure)

**Has realization**
This attribute is used to specify the process or activity that is the consequence of realization of the function.

For example,

- 282097004 | Ability to walk (observable entity) | 719722006 | Has realization (attribute) | of 870595007 | Walking (qualifier value)

**Inherent location**
This attribute is used to specify a body site or other location of the independent continuant in which the property exists.

For example,

- DNA taxon of Mycobacterium from bronchial secretions (observable entity) has 718497002 | Inherent location (attribute) | of bronchus

**Inheres in**
This attribute specifies the independent continuant in which the quality exists and on which the dependent quality (of this observable) depends.

For example,

- 307047009 | Core body temperature measured in rectum (observable entity) | has 704319004 | Inheres in (attribute) | of 278826002 | Body internal region (body structure)

**Precondition**
This attribute is used to specify body state, timing, challenges, or other situations that must be true of the entity to be observed.

For example,

- Plasma creatinine concentration 7 days post challenge (observable entity) has a Precondition of 7 days post challenge
163033001 |Lying blood pressure (observable entity)| has a 704326004 |Precondition (attribute)| of 102538003 |Recumbent body position (finding)|

**Procedure device**

This attribute is used to model devices associated with a procedure. This attribute is used to define high-level, general concepts that aggregate procedures according to the device involved.

**Process agent**

This attribute is used to specify the continuant (e.g. body structure or organism) that is causally active in the process on which the property depends. It is used to refine the meaning of the process named as the value of 704321009 |Characterizes (attribute)|, or it may simply repeat the meaning that is already there.

For example,

- Substance rate of secretion of somatotropin by pituitary following clonidine per os (observable entity) has the 704322002 |Process agent (attribute)| of 56329008 |Pituitary structure (body structure)|.

**Process duration**

This attribute specifies the duration of the process characterized by the observable property type.

For example,

- Mass rate of excretion of cortisone in 24 hour urine (observable entity) has the 704323007 |Process duration (attribute)| of 123027009 |24 hours (qualifier value)|

**Process extends to**

This attribute specifies that the process which the property characterizes has led to the inclusion of a previously not included structure.

For example,

The concept |Presence of direct invasion by primary malignant neoplasm of prostate to seminal vesicle (observable entity)| has a 1003703000 |Process extends to (attribute)| of |Seminal vesicle structure (body structure)|

**Process output**

This attribute is used to specify the substance or process produced by the process characterized by the observable property type.

For example,

- Substance rate of excretion of pregnanediol in micromoles per day (observable entity) has a 704324001 |Process output (attribute)| of 28268006 |Pregnanediol (substance)|
- 789350001 |Estimated quantity of intake of iron in 24 hours (observable entity)| has a 704324001 |Process output (attribute)| of iron

**Property**

This attribute is used to specify the type of feature (i.e. quality, disposition, function, or process characteristic) to be observed. Its values are abstract types of quality (length, odor, concentration) or abstract types of process features (rate, speed).

For example,

- Blood glucose mass concentration (observable entity) has the 370130000 |Property (attribute)| of 118539007 |Mass concentration (property) (qualifier value)|
Relative to
This attribute is used to specify the denominator of a relational property type, e.g. a ratio or proportion.

For example,
- Urine alpha aminobutyrate to creatinine ratio (observable entity) has 704325000 (Relative to (attribute)) 15373003 (Creatinine (substance))
- Neutrophils per 100 leukocytes in blood (observable entity) has 704325000 (Relative to (attribute)) 702962009 (Population of all leukocytes in portion of fluid (body structure))

Relative to part of
This attribute is used to specify the denominator of a relative relational property, such as a ratio of ratios.

For example,
- Relative substance concentration of cerebrospinal fluid IgM to plasma IgM (observable entity) has 719715003 (Relative to part of (attribute)) of 50863008 (Plasma (substance))

Scale Type
This attribute is used to specify the scale of the result of an observation or a diagnostic test (i.e., quantitative, qualitative, semi-quantitative).

- When defining observable entities for the international release, the Scale type (attribute) will not be used. Extensions are permitted to add specific subtypes of observable entities that include the Scale type (attribute), if desired.
- In instances where Observable entity content from SNOMED CT extensions that contain a SCALE TYPE relationship is promoted to the International release, the SCALE TYPE relationship will not be inactivated.

Technique
This attribute is used to specify the systematic method of an observation.

For example,
- Presence of Brucella abortus antibody in serum by latex agglutination (observable entity) has the 246501002 (Technique (attribute)) of 703448004 (Latex agglutination test technique (qualifier value))

Time Aspect
This attribute is used to specify the timing of an observation.

For example,
- Substance concentration of acetone in urine (observable entity) has the 370134009 (Time aspect (attribute)) of 123029007 (Single point in time (qualifier value))

Towards
This attribute is used to specify a disposition, what the disposition is towards, i.e. a specific triggering agent, or more generally, participant in the realization of the disposition.

For example,
- Quantitative susceptibility of Pseudomonas aeruginosa to amikacin in microbial isolate by disk diffusion (observable entity) has 704320005 (Towards (attribute)) of 387266001 (Amikacin (substance))
Units

This attribute is used to specify the units used in assigning a value to an observation.

For example,

- Basophils per 100 leukocytes (observable entity) has the 246514001 |Units (attribute)| of 415067009 |Percentage unit (qualifier value)|

Using device

This attribute is used to specify the instrument or equipment utilized to execute an action. Using device is appropriate when the device is actually used to carry out the action that is the focus of the procedure.

For example,

- 415921007 |Temperature of forehead using skin strip thermometer (observable entity)| has 424226004 |Using device (attribute)| of 448916003 |Skin strip thermometer (physical object)|

Observable Entity Naming Conventions

Test Observable Entity Naming Conventions

Naming conventions for the fully specified name (FSN) for observable entities and for naming evaluation procedures or observable entities that are submitted with names from the IFCC-IUPAC NPU systems are as follows:

General naming pattern: Property, Component, Direct Site

- First: Property
  - Property (the property type of the observable) is named first, when possible.
  - Modifier: Scale Method.
    - Scale Method refines the Property, and, therefore, precedes the action in the naming order.
      (Scale Method, Property)
    - Naming pattern: (Scale Method, Property), Component, Direct Site
  - Second: Component
    - Property is named first, followed by the entity that is the value of Component, when possible.
  - Third: Direct Site
    - Modifier: Time aspect. Time aspect provides information about the direct site and precedes it in the naming order. (Time aspect, Direct Site)

For example,

- 416125006 |Concentration of hemoglobin in erythrocyte (observable entity)|

Modeling: Screening measurements

Measurements done by screening should be specified with by screening method added at the end of the description.

Observable Entity Modeling

When observable entity concepts have been given a value, they behave like clinical findings, with respect to the concept model for context.

When observable entity concepts have not been given a value, they behave like procedures, with respect to the concept model for context.
Modeling

The observable entity model has been implemented in limited content areas in SNOMED CT thus far. Most of the content in the hierarchy is primitive.

- Over 165 observable entity concepts have been modeled describing physiological measurements (body temperature, respiratory rate, heart rate, blood pressure). Most of these are sufficiently defined using the attributes. [Vital sign (observable entity)] has been inactivated since it could not be universally defined.
- Additional concepts using observable entity attributes have been defined since the January 2020 release. The majority of the changes are related to nutritional intake (e.g., food intake, vitamin intake, fasting pattern) observable entity concepts.

Susceptibility observables should be modeled in accordance with the template specified here.

Observable Entity and Microbiology Test Results

When microbiology laboratory results are encoded, it is important to be aware of the context provided by the observation, i.e. the test performed and, therefore, the implied meaning of the result value, i.e. the organism.

For example, the combination of Logical Observation Identifiers Names and Codes (LOINC) for the lab test and SNOMED CT for the organism, provides a unique and specific meaning:

- LOINC provides microbiology reporting codes with attributes including the property through the use of PRID (presence or identity) and the scale through the use of NOM (nominal or categorical response that does not have a natural ordering) as the result value (typically the name of organism).
- Use of organism concepts in combination with such LOINC codes implies that a specific organism is seen, detected, identified, isolated, or present.

Organism

On its own, an organism concept can only indicate the definition of that organism. Its detection or presence can only be implied when it is paired with other information that may come from the electronic health application and/or from the LOINC observation.

Organism X or organism Y

Use organism X or organism Y when a laboratory report indicates a single isolate is assumed, but the lab is unable (for any reason) to differentiate the result instance.

For example,

- 703015006 | Human coxsackievirus or human echovirus (finding) |

Organism X, not organism Y

Use organism X, not organism Y when a laboratory report indicates a class of organisms described by the exclusion of specific Linnaean or non-Linnaean classes. These concepts are found in the organism hierarchy (based on reasonable use cases to avoid a combinatorial explosion). They are a primitive super class, in between the species or species subtype.

For example,

- 115407004 | Haemophilus influenzae, not b (organism) |

Genus X, not species Y and not species Z

Use Genus X, not species Y and not species Z when a laboratory report indicates a species of Genus X and confirms that it is not species Y, nor species Z. E.g. Bacillus species, not Bacillus anthracis and not Bacillus cereus (organism).

Use this naming convention only with Genus, species, and subspecies levels of the hierarchy.

Untypable organisms

Laboratory reports and journal articles may include an organism that could not be serotyped, e.g. E. coli, untypable. The requests for such concepts are declined due to ambiguity. Instead, use the closest taxonomic level in the hierarchy.
Presumptive values
Laboratory findings may be reported with a status of preliminary, presumptive, provisional, etc. These typically cover reportable or notifiable lab values. The status of a report is different from the result; it is part of the electronic health application model/message. The requests for such concepts are declined as they are ambiguous and subject to limitless combinations.

![Concepts with presumptive values](image)
Existing concepts with presumptive values are undergoing review for inactivation.

Mixed Organism
Some laboratories report findings indicating a mixed population of bacteria from several classes, e.g. *mixed anaerobic Gram negative bacilli*. The request for such a concept is added as a clinical finding. The actual organism is unknown, however there is a result, although more general.

Reporting Negative and Positive Results
Laboratories perform and report on specific tests to identify the absence, as well as the presence, of a particular pathogenic organism. Laboratories typically report negative result values, such as *X not seen*, *X not detected*, *X not isolated*, and *no X seen (or identified or isolated)* and positive results as *X seen*, *X detected*, and *X isolated*. The following tables includes the acceptable modeling for negative and positive results.

| Microbiology Tests: Reporting of Negative and Positive Values with Examples |
|--------------------|---------------------------------|---------------------------------------------------------------------------------|
| Lab test type (Observation) | Result value | Example lab test (e.g. SNOMED CT and/or LOINC term) | Example result value |
| General culture (where implied scale = nominal) | No X isolated (finding) | 61594008 [Microbial culture (procedure)] | Valid value 168204005 [Salmonella not isolated (finding)] |
| | X (organism) | 11475-1 [Microorganism identified in Unspecified specimen by Culture] | 27268008 [Genus Salmonella (organism)] |
| | | | Invalid value 264887000 [Not isolated (qualifier value)] |
| | | | 46651001 [Isolated (qualifier value)] |
| Organism Specific culture | Not isolated (qualifier value) | 122206002 [Bordetella pertussis culture (procedure)] | Valid value 264887000 [Not isolated (qualifier value)] |
| | Isolated (qualifier value) | 48741-3 [Bordetella pertussis; Nasopharynx; Culture] | 46651001 [Isolated (qualifier value)] |
| | | | Invalid value Bordetella pertussis not isolated Bordetella pertussis isolated |
### Microbiology Tests: Reporting of Negative and Positive Values with Examples

<table>
<thead>
<tr>
<th>General microscopic testing (where implied scale = Nominal)</th>
<th>No X seen (finding)</th>
<th>X (organism)</th>
<th>Procedure</th>
<th>Valid value</th>
<th>Invalid value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>609009000</td>
<td>Microscopic examination of bacterial smear of urine specimen (procedure)</td>
<td></td>
<td>27268008</td>
<td>Genus Salmonella (organism)</td>
</tr>
<tr>
<td></td>
<td>25145-4</td>
<td>Bacteria [Presence] in Urine sediment by Light microscopy</td>
<td></td>
<td>47492008</td>
<td>Not seen (qualifier value)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Specific microscopic testing</th>
<th>Not seen (qualifier value)</th>
<th>Present (qualifier value)</th>
<th>Procedure</th>
<th>Valid value</th>
<th>Invalid value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>408215009</td>
<td>Cryptosporidium microscopy (procedure)</td>
<td></td>
<td>47492008</td>
<td>Not seen (qualifier value)</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>52101004</td>
<td>Present (qualifier value)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Serologic, DNA or other organism specific test</th>
<th>Not detected (qualifier value)</th>
<th>Detected (qualifier value)</th>
<th>Procedure</th>
<th>Rationale: Almost all of these tests are organism-specific</th>
<th>Valid value</th>
<th>Invalid value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>871555000</td>
<td>Detection of ribonucleic acid of Severe acute respiratory syndrome coronavirus 2 (observable entity)</td>
<td></td>
<td>260415000</td>
<td>Not detected (qualifier value)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>95406-5</td>
<td>SARS-CoV-2 (COVID-19) RNA [Presence] in Nose by NAA with probe detection</td>
<td></td>
<td>260373001</td>
<td>Detected (qualifier value)</td>
<td></td>
</tr>
</tbody>
</table>

**Relationship between Observable Entities and NPU codes**

*Nomenclature, Properties, and Units (NPU)* is a coding system and terminology for identification and communication of examination results from clinical laboratories. Please see their website for more information: [NPU terminology](#).

Logically there is a relationship between NPU and SNOMED CT observable entities. A pilot project examined overlaps and possible alignment; further work may be done. However, there is no formally maintained SNOMED CT documentation on this alignment.

**Representing LOINC Terms with the SNOMED CT Observable Entity Model**

*Logical Observation Identifiers Names and Codes (LOINC)* terms are defined using the Observable Entity model in SNOMED CT as produced in the LOINC - SNOMED CT Cooperation Project releases.

The project release documentation contains information about how LOINC terms and parts are aligned with SNOMED CT concepts using the model.
Nutritional intake observable entities

Naming conventions for estimated and measured intake or nutrient administration, in aggregate or as a portion of intake via a specified route (i.e., oral, gastroenteral [enteral nutrition], parenteral nutrition, and via intravenous fluids), is as follows:


See the specific template here.

For example,

789106008 | Estimated quantity of intake of phosphorous in 24 hours (observable entity)
--- | ---
FSN: Estimated quantity of intake of phosphorous in 24 hours (observable entity)
PT: Estimated quantity of intake of phosphorous in 24 hours
Figure 1: Stated view of 789106008 |Estimated quantity of intake of phosphorous in 24 hours (observable entity)|

Dietary

"Dietary" is considered ambiguous and should not be included in SNOMED CT. Existing content that includes "dietary" will be considered for inactivation.

Observable Entity Templates

Templates are available for modeling quality observables and for process observables:

- Simple template for quality observables
- Simple template for process observables
  - Nutritional intake observables
- Susceptibility observables
  - The Susceptibility template is a disposition observable; there is no simple template at this time.

Neoplasm Observables

Observable entity concepts representing histopathology examination observables of neoplasms are being modeled according to a series of templates.

Please see the templates listed here: https://confluence.ihtsdotools.org/x/SonUAn. See the Community Content area for more information about the project.

Organism

<table>
<thead>
<tr>
<th>Definition</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Organisms of significance to human medicine</td>
<td>• 2265006 [Genus Candida (organism)]</td>
</tr>
<tr>
<td></td>
<td>• 710877000 [Beta lactam resistant bacteria (organism)]</td>
</tr>
</tbody>
</table>
Organism concepts
Organism concepts are used:
- In modeling cause of disease
- To document the cause of reportable or notifiable diseases
- In evidence-based infectious disease protocols, e.g. in clinical decision-support systems

Organisms with qualifiers

Intrinsic qualifiers
If a qualifier is an intrinsic part of an organism, it belongs in the organism hierarchy and is modeled accordingly. *Intrinsic* should be interpreted as a characteristic that is inherent in the organism (e.g. Gram-positive), as opposed to a context-dependent characteristic (e.g. some uses of *intracellular*).

When modeling organisms with qualifiers, the qualifier should be placed in front of the organism name.

Morphology qualifiers
For example, a non-Linnaean class of bacteria described by morphology
- 8745002 Gram-positive bacterium (organism)
- 416983001 Helical Gram-negative bacillus (organism)

Physiology qualifiers
For example, a non-Linnaean class of bacteria described by physiology
- 59343002 Anaerobic bacteria (organism)
- 417454003 Non-motile Salmonella (organism)

Resistance / susceptibility qualifiers
For example, A non-Linnaean class of bacteria described by antimicrobial susceptibility
- 712662001 Carbapenem resistant Enterobacteriaceae (organism)
- 417943000 Methicillin susceptible Staphylococcus aureus (organism)

Modeling with resistance-type qualifiers
Organisms with resistance-type qualifiers, i.e. where the qualifiers refer to the resistance phenotype and the organisms that are defined by the mechanism underlying the resistance phenotype, appear in the literature and are sometimes used interchangeably. However, in creating new concepts, these terms should be distinguished as they are separate concepts. For resistance-type qualifiers, use the antimicrobial agent as opposed to the enzyme that the organism is producing against the said antimicrobial agent.

For example,
- *Carbapenem resistant enterobacteriaceae* and *carbapenemase-producing enterobacteriaceae* share a significant overlap, but the former refers to the resistance phenotype, regardless of the mechanism of resistance. The presence of gene and carbapenemase production, as a resistance mechanism, usually results in clinically relevant levels of carbapenem resistance. However, it is possible to have only reduced susceptibility.

Validity
A number of qualifiers might be valid (e.g. aerobic microaerophilic, motile curved gram-negative bacteria). To determine the sequence, the decision-making process is stepwise as follows:
- Determined on a case-by-case basis
When requesting a new qualifier, an acceptable reference must be provided. Concepts with valid qualifiers are added to the International Release.

Organism groupings

Only authoritative taxonomic groupings are added to the SNOMED CT International Release. When requesting new organism concepts, authoritative references must be provided. Acceptance is determined on a case-by-case basis by authors. These concepts may evolve over time as the names evolve.

- Highly dependent on fitting in with the model limitations
- Based on Bergey’s Manual of Systematic Bacteriology as the primary reference

**Complex or Group**

The terms “complex” and “group” are often used in scientific papers. Laboratories then reflect the words they see in those papers in their local descriptions. However, the terms used in scientific papers are not authoritative taxonomic groupings; rather, they are just concepts used for ease of publication and grouping sets of organisms that are similar in certain functions or structure.

Implementers must be aware these types of concepts may evolve over time. As the sophistication of microbiology labs increases, the “members” of each complex may change and the complex concepts actually become obsolete. For example, this has occurred for some of the Centers for Disease Control and Prevention (CDC) groups where a number of these concepts have actually been given names and the CDC group name is archaic.

When requesting a new group or complex, an acceptable authoritative reference must be provided. The reference should clearly specify the list of species and subspecies associated with the complex/group.

Existing complex or group concepts, with grouper concepts separate from the genus, but with the same meaning as the genus, will be inactivated in the SNOMED CT International Release.

Descriptions with group or complex as synonyms of the genus, will be deprecated from the SNOMED CT International Release (The genus concept should be used for these concepts).

**Microorganisms**

Microorganism is a common grouping name for organisms, but it does not align with Linnaean classification. Microorganisms are organisms that can only be seen using microscopy. Four major classes could reasonably be assigned to microorganism at the highest levels. Viruses, prions, bacteria and archaea are all microscopic. Fungi are both microscopic and macroscopic and this is also true for animals. Finally, there are examples of organisms (e.g. Phylum Nemata) that are macroscopic as adults but diagnostic life-cycle stages such as eggs and larvae are microscopic. Assigning and maintaining all subtypes to this seemingly familiar organism group is problematic and would be time and resource intensive. This concept has been deprecated and will not be added to the organism hierarchy.

Biotype, Serotype, Serogroup

Requests for new concepts are evaluated on a case-by-case basis.

It is important to understand the meaning from the requestor and determine how it can be modeled.

These concepts may evolve over time as the names evolve.

**X-like Organism**

"X-like" organism is a term construction used in the medical lexicon that is outside the classic Linnaean taxonomy. "X-like" organisms are identified by their similarity to some other organism. There is no single category or use of X-like organism terms; the meaning of these terms is context-dependent and open to interpretation when no
context is provided. For many of these terms, the meaning will change with time. In some cases, this leads to a chain of terms that remains in colloquial use but loses value and place in the scientific literature. In addition, while reporting X-like organisms is clinically significant—unlike “untypable” concepts—they cannot have a specific parent. These concepts are added:

- only if clear context is provided by the requester; and
- under the highest level concepts in the “organism” hierarchy i.e. direct parents would be Virus, Bacteria, Fungus.

Provisional serotypes

Provisional serotypes, i.e. serotypes that have been defined but not given a number in the antigenic schema, are considered for addition on an ad hoc basis and only if it can be confirmed that this is a reproducible assignment not being duplicated by multiple organizations.

Multidrug-resistant, extensively drug-resistant, pan drug-resistant bacteria

SNOMED International adopted the recommendations of a joint initiative of the European Centre for Disease Prevention and Control (ECDC) and the CDC for the characterization of the different patterns of resistance found in healthcare-associated, antimicrobial resistant bacteria. A panel of international experts convened and drafted a proposal which provides clear consensus definitions. Please refer to the following article for details: Magiorakos, A. Srinivasan, A. Multidrug-resistant, extensively drug-resistant and pandrug-resistant bacteria: an international expert proposal for interim standard definitions for acquired resistance. Clinical Microbiol Infect 2012; 18: 268-281.

Organism Naming Conventions

Fully Specified Name

The fully specified name (FSN) of organism concepts, names classes that are officially recognized Linnaean taxonomic classes (other than species), and include a designation of rank. They include, but are not limited to Phylum, Order, Suborder, Class, Family, Genus, and subspecies.

Properly constructed FSNs contain terms indicating the taxonomic rank + the recognized name of that rank + semantic tag.

For example,

- 106544002 | Family Enterobacteriaceae (organism)

Rank

The naming convention is not applied to concepts that only refer to a subgroup of a rank.

For example,

- Vancomycin resistant Enterococcus (organism) is correct in capitalizing Enterococcus. It refers to a subclass of the genus, Enterococcus species that are resistant. Enterococcus is a scientific name of an organism class; therefore, the first letter is capitalized.
- Vancomycin resistant Genus Enterococcus is incorrect. It refers to the rank only, Genus Enterococcus.
Official names of organisms may include abbreviations such as “subsp.” and “subgen.” (Domain Bacteria and Kingdom Plantae). Official names of organisms may also include parentheses e.g. “Cypraea (Cypraea) tigris” (Kingdom Animalia) and “Bacillus (subgen. Bacillus Cohn 1872, 174) subtilis” (Domain Bacteria).

- The FSN of organisms should include the expanded word for rank i.e. “subgenus” or “subspecies” and not an abbreviation of same.
- The FSN should not include parentheses.

For example

- Genus Pleione subgenus Scopulorum (organism)
- Genus Cypraea subgenus Cypraea tigris (organism)
- Staphylococcus succinus subspecies casei (organism)

Preferred Term

The Preferred Term is the official scientific name. It may include abbreviations and/or parentheses. The preferred term usually does not include the taxonomic rank designation except for the following cases:

- Official names of organisms may include abbreviated taxonomic rank such as “subg.” “subgen.” or “subsp.”.
- In rare cases, two Linnaean taxon ranks in the same hierarchy may have the same name. For example, in Kingdom Bacteria, “Thermodesulfobacteria” is an applicable term at both Phylum and Class levels. In this case, the taxon rank is included in the preferred term to prevent any ambiguity.

For example,

- Cypraea (Cypraea) tigris
- Pleione subg. Scopulorum
- Bacillus (subgen. Bacillus Cohn 1872, 174) subtilis
- Staphylococcus succinus subsp. casei
Qualifiers in organism names

When modeling organisms with qualifiers, the qualifier should be placed in front of the organism name.

Organism class variants

The description of organism classes that are subspecies subtypes and variants may include terms such as serogroup, serotype, biotype, variant, biovar, serovar, and pathovar.

For example,

- 698206009 |Brucella suis biovar 4 (organism)|

The subspecies types and variants should be included in the FSN, PT and other descriptions. This is to avoid ambiguity when the same number or letter is used to refer to different organism variants.

For example, without mentioning the specific variant (serogroup vs. serotype) and the nomenclature system (Danish vs. American), "Streptococcus pneumoniae 48" can refer to the following:

- Streptococcus pneumoniae Danish serotype 48 (which is equivalent to Streptococcus pneumoniae American serotype 82)
• Streptococcus pneumoniae American serotype 48 (which is equivalent to Streptococcus pneumoniae Danish serotype 7B)
• Streptococcus pneumoniae serogroup 48

Abbreviations (var, var., sv, sv., bv, bv., pv, pv.) must not be used in the FSN.

Capitalization of organism names and binomial format

Official scientific names for organisms should be capitalized. The designation of rank does not require capitalization.

For example,

• 426813007 | Order Acidobacteriales (organism) | has case significance of “Initial character case insensitive”

There is an exception to the above guidelines where the binomial format for an organism species includes capitalization of the genus name but the species name begins with a lower-case letter.

For example,

• 24224000 | Brucella abortus (organism) |

Salmonella serotype nomenclature

Salmonella serotypes have a quadrinomial format of Genus species subspecies Serotype where the serotype name is capitalized.

For example,

• A synonym for 114683003 | Salmonella Doel (organism) | is Salmonella enterica subsp. enterica ser. Doel

Additional descriptions, without the species and subspecies names, are in common usage for Salmonella serotypes.

For example,

• 656008 | Salmonella Os (organism) |

In SNOMED CT, the serotype name in the description should be capitalized.

⚠️ Salmonella Serotypes

Salmonella serotypes, without the species and subspecies names, should not be confused with binomial species names of other organisms.

Streptococcus pneumoniae

Streptococcus pneumoniae is a human pathogen whose virulence is based on its protective polysaccharide capsule. Study of the polysaccharide capsule has identified multiple serogroups and serotypes. Serotypes are defined by the chemical structure and immunologic properties of their polysaccharide; each serogroup contains one or more serotypes that elicit the same antibody response.

There are two serotype naming systems, one in the U.S. and one in Denmark. The Danish system is nearly universally accepted and preferred. For details, please refer to See Geno K A, Gilbert G L, Song J Y, Skovsted I C, Klugman K P, Jones C, Konradsen H B, Nahm M H. Pneumococcal capsules and their types: past, present, and future. Clinical Microbiology Reviews 2015; 28(3):871-899. [PMID: 26085553]).

Streptococcus pneumoniae concepts

A review of Streptococcus pneumoniae serotypes in SNOMED CT showed lack of specificity, as well as inconsistency, in the naming of Streptococcus pneumoniae serotypes. Guidelines for creating concepts containing Streptococcus pneumoniae serotypes were formulated. They are as follows:
FSN and preferred term (PT) descriptions should follow the Danish naming system. When an American synonym exists, it should be added. A synonym (SYN) that matches the FSN, but does not contain the naming system can also be added.

For example,

698149000 | Streptococcus pneumoniae serotype 48 (organism) |

- FSN: Streptococcus pneumoniae Danish serotype 48 (organism)
- PT: Streptococcus pneumoniae Danish serotype 48
- SYN: Streptococcus pneumoniae American serotype 82
- SYN: Streptococcus pneumoniae serotype 48

The guidelines for creating new concepts containing Streptococcus pneumoniae serotypes also apply to concepts in other SNOMED CT hierarchies, such substances and procedures.

For example,

120683007 | Streptococcus pneumoniae serotype 7F antibody (substance) |

- FSN: Antibody to Streptococcus pneumoniae Danish serotype 7F (substance)
- PT: Streptococcus pneumoniae Danish serotype 7F Ab
- SYN: Antibody to Streptococcus pneumoniae Danish serotype 7F
- SYN: Anti-Streptococcus pneumoniae Danish serotype 7F antibody
- SYN: Streptococcus pneumoniae Danish serotype 7F antibody
- SYN: Antibody to Streptococcus pneumoniae American serotype 51

Legacy Streptococcus pneumoniae concepts
FSNs that adhered to one of the naming systems were kept, but changes were made to the descriptions, based on the above guidelines. Any resulting duplicates were deprecated.

FSNs that did not adhere to one of the naming system were inactivated as ambiguous. They were replaced with newly created concepts, based on the above guidelines.

Missing serotype concepts were added.

Influenza virus nomenclature
Follow the latest names for genus and species according to the taxonomy authority. Although the genus and species names for influenza viruses are similar, they each follow a distinct pattern, which should be used in SNOMED CT. Also, the name of the virus should always be capitalized.

For species, the word virus is included as a separate word and follows the letter designation.

For example,

- 407482004 | Influenza C virus (organism) |
- 710661004 | Immunoglobulin M antibody to Influenza B virus (substance) |
- 1067491100119108 | Otitis media caused by Influenza A virus (disorder) |

For genus, virus is included in the genus name and is not a separate word.

For example,

- 407481006 | Genus Gammainfluenzavirus (organism) |
- 407477006 | Genus Alphainfluenzavirus (organism) |

The disorder influenza need not be capitalized.

For example,

- 16311000119108 | Pneumonia caused by influenza (disorder) |
- 309789002 | Encephalitis caused by influenza (disorder) |
US/GB spelling variants for taxonomic concepts

Taxonomic resources (e.g. Integrated Taxonomic Information System or ITIS, List of Prokaryotic names with Standing in Nomenclature or LPSN) use the official scientific name for organisms. Similarly, in SNOMED CT, the official scientific name should be used in FSNs and PTs. For descriptions representing common names, if the spelling in a country or region is different, the preferred spelling should be added in the language RefSet extension as a synonym.

Use of X species

In the context of the Linnaean organism hierarchy, there is no difference between Salmonella species and simply Salmonella, the genus. Terms with X species, such as Salmonella species, are routinely used in laboratory reporting. They may provide additional information, other than the place of the organism in the Linnaean hierarchy. However, the intended connotation may vary from lab to lab and from organism to organism.

Since the organism concept represents a class of organisms, it cannot also represent what was, was not, or what will be done to identify the organism. Neither can it represent other information about the result. If there is additional information to report, it should be in a separate statement or comment (e.g. further species identification pending or sent to reference laboratory for further identification or further identification to be done if clinically indicated).

**X species**

Addition of X species as a description to genus X is allowed and is done per request.

Microorganism name changes

Microorganism taxonomic names may change, often due to scientific advances. This may result in:

- Finding an organism in a particular taxonomic group (e.g. Genus) that is unrelated, on a molecular basis, to other members of the group.
- Reassessing the taxonomic group originally established, based on phenotypic characteristics.
- Proposing to reassign the organism to a different existing or new taxonomic group.

On a case by case basis, requests for name changes are based on the following use cases:

The name of an organism changes. This scenario is also applicable when an organism name changes on multiple occasions over time.

- Change the FSN for affected concepts, but not the concept ID, by creating a new FSN and description. Inactivate the old FSN with an inactivation value of Outdated.
- Retain the old name as a synonym.

A single species is reclassified as multiple species. This scenario is applicable if the change in classification happens at a single point in time and is reflected as such in the authoritative resources.

- Create the new concepts.
- Inactivate the original concept as ambiguous.
- Set a possibly equivalent to relationship between the old concept and the new concepts.

Multiple species are reclassified as one. This scenario is applicable if the change in classification happens at a single point in time and is reflected as such in the authoritative resources.

- Create a new concept.
- Inactivate the existing concepts as outdated with replaced by relationships to the new concept.
Organism life stages

Concepts in the organism hierarchy represent fully realized organisms. An organism’s life cycle stage is a characteristic of a given taxon. It represents different stages of life e.g. egg, larva, and adult.

Organism stages themselves are characteristics common to members of a given taxon.

SNOMED CT allows for the representation of an organism in a specific life cycle stage.

For example,
- 337915000 | Homo sapiens (organism) | are organisms. Homo sapiens include humans, in general, as well as children.
- Childhood is a life cycle stage, however it is not an organism.

Similarly,
- An egg of a particular nematode, e.g. 42625000 | Strongyloides stercoralis (organism) | is an organism. It is alive and can pass through other stages appropriate to its species.

However, the egg stage of Strongyloides stercoralis is not an organism. Many diagnostic test results, identify organisms ‘participating’ in particular life cycle stages.

For example, the results of a 83033005 | Fecal analysis (procedure) | may identify the presence of 609326000 | Larva of Strongyloides stercoralis (organism) | and 699572004 | Egg of Strongyloides stercoralis (organism) |

Organism concepts

Concepts in the organism hierarchy should not represent organism structures (e.g. fungal hyphae). In addition, the word "stage" should be excluded from concepts representing life cycle of an organism (e.g. larval stage of a nematode parasite). This does not preclude representations of organisms ‘participating’ in a specific stage of life e.g. 609061000 | Larva of genus Ascaris (organism) |

Naming patterns

FSN pattern: (Life cycle stage) of (Taxon including rank, if required) (organism)

For example,
- 609043009 | Adult of phylum Nemata (organism) |
- 699572004 | Egg of Strongyloides stercoralis (organism) |

The name of the rank is included with the first letter lower case, except at the species and subspecies levels, where the Linnaean binomial and trinomial are specified.

PT pattern: (Taxon including rank, if required) (life cycle stage)

For example,
- Phylum Nemata adult
- Strongyloides stercoralis egg

Cestode larvae

A number of cestode larvae have historically been referred to using Linnaean binomial names that are completely different from corresponding adult (or egg) names.

For example,
- 47399003 | Larva of Taenia saginata (organism) |, a human tapeworm, is usually called Cysticercus bovis.

PT pattern: Linnaean binomial of larva OR (Taxon including rank if required) (life cycle stage)

For example,
• Cysticercus bovis
• Cysticercus cellulosae
• Class Cestoda larva

Although rare, a subtype of cestode larva may appear to be a Linnaean trinomial name. This, then, is the PT:
• Diphyllobothrium latum sparganum

Other acceptable synonyms
Some organisms and stages are referred to in an *adjectival* form (e.g. Ascarid egg) or by common name (e.g. adult nematode). When used (especially when described as part of a request), these terms may be included as additional synonyms.

Resources for organism naming
SNOMED International utilizes various resources when reviewing changes to the organism hierarchy. They include:

**Bacteria**
- List of Prokaryotic names with Standing in Nomenclature (LPSN)
- International Committee on Systematics of Prokaryotes (ICSP)
- International Journal of Systematic and Evolutionary Microbiology
- DSMZ-Prokaryotic Nomenclature Up-to-date

**Fungi**
- MycoBank Database
- Index Fungorum

**Viruses**
- International Committee on Taxonomy of Viruses (ICTV)

**Parasites**
- National Center for Biotechnology Information (NCBI) Taxonomy (Although not an authoritative source, provides useful links to other sources; used by Unified Medical Language System (UMLS) as a QA source)

**General**
- Integrated Taxonomic Information System (ITIS) (Covers a limited number of organisms and is not up-to-date on all areas. For bacteria, fungus, and virus, consult resources noted above as primary references.)

**Pharmaceutical and Biologic Product**
The Pharmaceutical / biologic product hierarchy is comprised of multiple smaller hierarchies, e.g. the Medicinal product hierarchy.

Editorial guidelines for the Pharmaceutical / biologic product hierarchy are available in the Pharmaceutical/Biologic Product Editorial Guide.

**Pharmaceutical and Biologic Product Attributes Summary**
Editorial guidelines for the 373873005 |Pharmaceutical / biologic product (product)| hierarchy are available in the Pharmaceutical and Biologic Product Hierarchy SNOMED CT Editorial Guide.

When authoring in this domain, these are the approved attributes and allowable ranges. They are the Human Readable Concept Model (HRCM).
### Domain Information for 373873005 Pharmaceutical / biologic product (product)

| Domain Constraint | << 373873005 Pharmaceutical / biologic product (product) |
| Parent Domain     | - |
| Proximal Primitive Constraint | << 373873005 Pharmaceutical / biologic product (product) |
| Proximal Primitive Refinement | - |

### Author View of Attributes and Ranges for 373873005 Pharmaceutical / biologic product (product)

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Grou ped</th>
<th>Cardi nality</th>
<th>In Grou p Cardi nality</th>
<th>Range Constraint</th>
</tr>
</thead>
<tbody>
<tr>
<td>1142140007</td>
<td>0..1</td>
<td>0..0</td>
<td>int(&gt;#0..)</td>
<td></td>
</tr>
<tr>
<td>1142140006</td>
<td>0..1</td>
<td>0..0</td>
<td>int(&gt;#0..)</td>
<td></td>
</tr>
<tr>
<td>1142139005</td>
<td>0..1</td>
<td>0..0</td>
<td>int(&gt;#0..)</td>
<td></td>
</tr>
<tr>
<td>127489000</td>
<td>0..1</td>
<td>0..1</td>
<td>&lt;&lt; 105590001 Substance (substance)</td>
<td></td>
</tr>
<tr>
<td>732943007</td>
<td>0..1</td>
<td>0..1</td>
<td>&lt;&lt; 105590001 Substance (substance)</td>
<td></td>
</tr>
<tr>
<td>73722007</td>
<td>0..1</td>
<td>0..1</td>
<td>767524001 Unit of measure (qualifier value)</td>
<td></td>
</tr>
<tr>
<td>1142137007</td>
<td>0..*</td>
<td>0..1</td>
<td>dec(&gt;#0..)</td>
<td></td>
</tr>
<tr>
<td>733725009</td>
<td>0..*</td>
<td>0..1</td>
<td>767524001 Unit of measure (qualifier value)</td>
<td></td>
</tr>
<tr>
<td>1142138002</td>
<td>0..*</td>
<td>0..1</td>
<td>dec(&gt;#0..)</td>
<td></td>
</tr>
<tr>
<td>762951001</td>
<td>0..*</td>
<td>0..1</td>
<td>&lt;&lt; 105590001 Substance (substance)</td>
<td></td>
</tr>
<tr>
<td>860779006</td>
<td>0..*</td>
<td>0..*</td>
<td>&lt;&lt; 362981000 Qualifier value (qualifier value)</td>
<td></td>
</tr>
<tr>
<td>1149366004</td>
<td>0..*</td>
<td>0..1</td>
<td>1149484003 Ingredient qualitative strength (qualifier value)</td>
<td></td>
</tr>
<tr>
<td>411116001</td>
<td>0..1</td>
<td>0..0</td>
<td>&lt;&lt; 736542009 Pharmaceutical dose form (dose form)</td>
<td></td>
</tr>
<tr>
<td>762949000</td>
<td>0..*</td>
<td>0..1</td>
<td>&lt;&lt; 105590001 Substance (substance)</td>
<td></td>
</tr>
<tr>
<td>732947000</td>
<td>0..*</td>
<td>0..1</td>
<td>&lt;&lt; 767524001 Unit of measure (qualifier value)</td>
<td></td>
</tr>
<tr>
<td>1142136003</td>
<td>0..*</td>
<td>0..1</td>
<td>dec(&gt;#0..)</td>
<td></td>
</tr>
<tr>
<td>732945000</td>
<td>0..*</td>
<td>0..1</td>
<td>767524001 Unit of measure (qualifier value)</td>
<td></td>
</tr>
<tr>
<td>860781008</td>
<td>0..*</td>
<td>0..0</td>
<td>&lt;&lt; 362981000 Qualifier value (qualifier value)</td>
<td></td>
</tr>
<tr>
<td>774158006</td>
<td>0..1</td>
<td>0..0</td>
<td>&lt;&lt; 774167006 Product name (product name)</td>
<td></td>
</tr>
<tr>
<td>774159003</td>
<td>0..1</td>
<td>0..0</td>
<td>&lt;&lt; 774164004 Supplier (supplier)</td>
<td></td>
</tr>
<tr>
<td>Code</td>
<td>Description</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>----------------------</td>
<td>-----------------------------------------------------------------------------</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1149367008</td>
<td>Has target population (attribute)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>7630320000</td>
<td>Has unit of presentation (attribute)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>7669390001</td>
<td>Plays role (attribute)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>11487930005</td>
<td>Unit of presentation size quantity (attribute)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>120091000221107</td>
<td>Unit of presentation size unit (attribute)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>781405001</td>
<td>Medicinal product package (product)</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>736542009</td>
<td>Pharmaceutical dose form (dose form)</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Domain Information for 781405001 Medicinal product package (product)**

- **Domain Constraint**: << 781405001 Medicinal product package (product)
- **Parent Domain**: 373873005 Pharmaceutical / biologic product (product)
- **Proximal Primitive Constraint**: << 781405001 Medicinal product package (product)
- **Proximal Primitive Refinement**: -

**Author View of Attributes and Ranges for 781405001 Medicinal product package (product)**

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Grouped</th>
<th>Cardinality</th>
<th>In Group Cardinality</th>
<th>Range Constraint</th>
</tr>
</thead>
<tbody>
<tr>
<td>774160008 Contains clinical drug</td>
<td>1</td>
<td>1..*</td>
<td>1..1</td>
<td>&lt;&lt; 763158003 Medicinal product (product)</td>
</tr>
<tr>
<td>1142143009 Count of clinical drug</td>
<td>0</td>
<td>1..1</td>
<td>0..0</td>
<td>int(&gt;=0..)</td>
</tr>
<tr>
<td>1142142004 Has pack size</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
<td>int(&gt;=0..)</td>
</tr>
<tr>
<td>774163005 Has pack size unit</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
<td>&lt;&lt; 767524001 Unit of measure (qualifier value)</td>
</tr>
</tbody>
</table>

**Domain Information for 736542009 Pharmaceutical dose form (dose form)**

- **Domain Constraint**: << 736542009 Pharmaceutical dose form (dose form)
- **Parent Domain**: -
- **Proximal Primitive Constraint**: << 736542009 Pharmaceutical dose form (dose form)
- **Proximal Primitive Refinement**: -

**Author View of Attributes and Ranges for 736542009 Pharmaceutical dose form (dose form)**

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Grouped</th>
<th>Cardinality</th>
<th>In Group Cardinality</th>
<th>Range Constraint</th>
</tr>
</thead>
</table>

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## Domain Information for 736478001 [Basic dose form (basic dose form)]

**Domain Constraint**

<< 736478001 [Basic dose form (basic dose form)]

**Parent Domain**

- 

**Proximal Primitive Constraint**

<< 736478001 [Basic dose form (basic dose form)]

**Proximal Primitive Refinement**

- 

## Author View of Attributes and Ranges for 736478001 [Basic dose form (basic dose form)]

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Grouped</th>
<th>Cardinality</th>
<th>In Group Cardinality</th>
<th>Range Constraint</th>
</tr>
</thead>
<tbody>
<tr>
<td>736518005  Has state of matter (attribute)</td>
<td>0</td>
<td>1..1</td>
<td>0..0</td>
<td>&lt; 736471007 [State of matter (state of matter)]</td>
</tr>
</tbody>
</table>

### Physical Force

**Definition**

Forces applied to the body that may cause injury

**Examples**

- 87588000 [High altitude (physical force)]
- 263762005 [Friction (physical force)]

⚠️ **Physical Force**

The concepts in the Physical force hierarchy primarily represent physical forces that may play a role in injuries.

### Physical Object

**Definition**

Physical devices relevant to healthcare or to injuries/accidents

**Examples**

- 469785004 [Heel protector (physical object)]
- 40388003 [Implant, device (physical object)]
Editorial guidelines for the Physical object hierarchy are available on the Devices Project page here: Devices Project.

Use case
Concepts in the Physical object hierarchy include natural and man-made objects. One use for these concepts is modeling procedures that use devices (e.g. catheterization).

Scope
In scope
Concepts representing hardware or software...

Out of scope
IVD, prescribing status...

Physical Object Attributes Summary
When authoring in this domain, these are the approved attributes and allowable ranges. They are from the Human Readable Concept Model (HRCM).

| Domain Information for 260787004 | Physical object (physical object) |
| Domain Constraint | << 260787004 | Physical object (physical object) |
| Parent Domain | - |
| Proximal Primitive Constraint | << 260787004 | Physical object (physical object) |
| Proximal Primitive Refinement | - |
Physical Object Defining Attributes

The following defining attributes correspond to the Physical Object Attributes Summary table.

Has compositional material

This attribute represents the material or substance of which an object is composed.

For example,
- 12345 |x (physical object) |

Has device characteristic

This attribute represents a device characteristic such as how it is expected to react with the body or whether it has been prepared specifically for a subject.

For example,
- 12345 |x (physical object) |

Has device intended site

This attribute represents the site where the device is intended to reside in or on the body.

For example,
- 12345 |x (physical object) |

Has filling

This attribute represents the material or substance that fills an object.

For example,
• 12345 | x (physical object) | ...

Has surface characteristic
This attribute represents the tactile characteristic and appearance of the surface of an object.

For example,
• 12345 | x (physical object) | ...

Physical Object Naming and Modeling Conventions
Specific editorial guidelines for modeling and terming will be documented for each device type as completed.

Subpage ex Breast prostheses
Subpage ex Cardiac stents
Subpage ex Catheters

Procedure

<table>
<thead>
<tr>
<th>Definition</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Procedure: activities performed in the provision of health care (includes medical history-taking, physical examination, diagnostic and therapeutic interventions, training and education, and counseling)</td>
<td>• 54321008</td>
</tr>
<tr>
<td>• Regime/therapy (subtype of procedure): set of procedures focused on a single purpose on one patient over time (e.g. repeated administration of drug in a small dose for an indefinite period of time)</td>
<td>• 386513007</td>
</tr>
</tbody>
</table>

Procedure concepts
Procedure concepts represent activities performed in the provision of health care. This hierarchy represents a broad variety of activities, including but not limited to:

• Invasive procedures, e.g. 77018005 | Excision of lesion of intracranial artery (procedure) |
• Administration of medicines, e.g. 39343008 | Pertussis vaccination (procedure) |
• Imaging procedures, e.g. 47079000 | Ultrasonography of breast (procedure) |
• Education procedures, e.g. 183063000 | Low salt diet education (procedure) |
• Administrative procedures, e.g. 305212007 | Medical records transfer (procedure) |

Procedure Attributes Summary
When authoring in this domain, these are the approved attributes and allowable ranges. They are from the Human Readable Concept Model (HRCM). In addition, 386053000 | Evaluation procedure (procedure), 387713003 | Surgical procedure (procedure), and 433590000 | Administration of substance via specific route (procedure) each have unique defining attributes as seen in their separate tables below.

Domain Information for 71388002 | Procedure (procedure) |
Domain Constraint | <= 71388002 | Procedure (procedure) |
Parent Domain | -
## Proximal Primitive Constraint

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Grouped/Cardinality</th>
<th>Range Constraint</th>
</tr>
</thead>
</table>
| Access (attribute) | 1 0..* 0..1 | << 309795001 | Surgical access values (qualifier value) |}
| Direct device (attribute) | 1 0..* 0..1 | << 49062001 | Device (physical object) |}
| Direct morphology (attribute) | 1 0..* 0..1 | << 49755003 | Morphologically abnormal structure (morphologic abnormality) |}
| Direct substance (attribute) | 1 0..* 0..1 | << 105590001 | Substance (substance) OR << 373873005 | Pharmaceutical / biologic product (product) |}
| Has focus (attribute) | 1 0..* 0..1 | << 40464003 | Clinical finding (finding) OR << 71388002 | Procedure (procedure) |}
| Has intent (attribute) | 1 0..* 0..1 | << 363675004 | Intent (nature of procedure values) (qualifier value) |}
| Indirect device (attribute) | 1 0..* 0..1 | << 49062001 | Device (physical object) |}
| Indirect morphology (attribute) | 1 0..* 0..1 | << 49755003 | Morphologically abnormal structure (morphologic abnormality) |}
| Method (attribute) | 1 0..* 0..1 | << 129264002 | Action (qualifier value) |}
| Priority (attribute) | 1 0..* 0..1 | << 272125009 | Priorities (qualifier value) |}
| Procedure device (attribute) | 1 0..* 0..* | << 49062001 | Device (physical object) |}
| Procedure morphology (attribute) | 1 0..* 0..* | << 49755003 | Morphologically abnormal structure (morphologic abnormality) |}
| Procedure site (attribute) | 1 0..* 0..* | << 442083009 | Anatomical or acquired body structure (body structure) |}
| Procedure site - Direct (attribute) | 1 0..* 0..1 | << 442083009 | Anatomical or acquired body structure (body structure) |}
| Procedure site - Indirect (attribute) | 1 0..* 0..1 | << 442083009 | Anatomical or acquired body structure (body structure) |}
| Recipient category (attribute) | 1 0..* 0..1 | << 125676002 | Person (person) OR << 133928008 | Community (social concept) OR << 33539004 | Family (social concept) OR << 38919008 | Group (social concept) |}
| Revision status (attribute) | 1 0..* 0..1 | << 255231005 | Revision - value (qualifier value) OR << 257958009 | Part of multistage procedure (qualifier value) OR << 261424001 | Primary operation (qualifier value) |}
| Using access device (attribute) | 1 0..* 0..1 | << 49062001 | Device (physical object) |}
| Using device (attribute) | 1 0..* 0..* | << 49062001 | Device (physical object) |}
| Using energy (attribute) | 1 0..* 0..1 | << 78621006 | Physical force (physical force) |}
| Using substance (attribute) | 1 0..* 0..1 | << 105590001 | Substance (substance) |}

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### Domain Information for 386053000 Evaluation procedure (procedure)

**Domain Constraint**
- `< 386053000 Evaluation procedure (procedure)`

**Parent Domain**
- 71388002 Procedure (procedure)

**Proximal Primitive Constraint**
- `< 71388002 Procedure (procedure)`

**Proximal Primitive Refinement**
- `[1..*] 260686004 Method = [[+id(<< 129265001 Evaluation - action)]]`

### Author View of Attributes and Ranges for 386053000 Evaluation procedure (procedure)

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Grouped</th>
<th>Cardinality</th>
<th>In Group Cardinality</th>
<th>Range Constraint</th>
</tr>
</thead>
<tbody>
<tr>
<td>246093002 Component (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
<td><code>&lt; 105590001 Substance (substance) OR &lt; 123037004 Body structure (body structure) OR &lt; 123038009 Specimen (specimen) OR &lt; 260787004 Physical object (physical object) OR &lt; 373873005 Pharmaceutical / biologic product (product) OR &lt; 41067006 Organism (organism) OR &lt; 419891008 Record artifact (record artifact)</code></td>
</tr>
<tr>
<td>116686009 Has specimen (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
<td><code>&lt; 123038009 Specimen (specimen)</code></td>
</tr>
<tr>
<td>370129005 Measurement method (attribute)</td>
<td>1</td>
<td>0..*</td>
<td>0..1</td>
<td><code>&lt; 127789004 Laboratory procedure categorized by method (procedure)</code></td>
</tr>
<tr>
<td>370130000 Property (attribute)</td>
<td>1</td>
<td>0..1</td>
<td>0..1</td>
<td><code>&lt; 118598001 Property (qualifier value)</code></td>
</tr>
<tr>
<td>370132008 Scale type (attribute)</td>
<td>1</td>
<td>0..1</td>
<td>0..1</td>
<td><code>&lt; 117362005 Nominal value (qualifier value) OR &lt; 117363000 Ordinal value (qualifier value) OR &lt; 117364006 Narrative value (qualifier value) OR &lt; 117365007 Ordinal OR quantitative value (qualifier value) OR &lt; 117444000 Text value (qualifier value) OR &lt; 26716007 Qualitative (qualifier value) OR &lt; 30766002 Quantitative (qualifier value)</code></td>
</tr>
<tr>
<td>370134009 Time aspect (attribute)</td>
<td>1</td>
<td>0..1</td>
<td>0..1</td>
<td><code>&lt; 7389001 Time frame (qualifier value)</code></td>
</tr>
</tbody>
</table>

### Domain Information for 387713003 Surgical procedure (procedure)

**Domain Constraint**
- `< 387713003 Surgical procedure (procedure)`

**Parent Domain**
- 71388002 Procedure (procedure)

**Proximal Primitive Constraint**
- `< 71388002 Procedure (procedure)`

**Proximal Primitive Refinement**
- `[1..*] 260686004 Method = [[+id(<< 129284003 Surgical action (qualifier value)]]`
Procedure Defining Attributes

The following contain the defining attributes for Procedure concepts. Evaluation, Surgical, and Administration of Substance via Specific Route procedures each have unique defining attributes.

Procedure, General

The following defining attributes correspond to the Procedure Attributes Summary table from the HRCM.

Self-grouped Attributes

The following attributes are self-grouped, meaning they are not grouped with any other attributes:

- Priority
- Has focus

Access

Access (attribute) describes the route used to access the site of a procedure. It distinguishes open, closed, and percutaneous procedures.

For example,

- 174572001 Open removal of bile duct stent (procedure) has Access of Open approach - access (qualifier value)

Direct device

Direct device (attribute) represents the device on which the method directly acts.
For example,

- **431698006** | Adjustment of gastric banding using fluoroscopic guidance (procedure) | has Direct device of Surgical band (physical object)

**Direct morphology**

Direct morphology (attribute) describes the morphologically abnormal structure that is the direct object of the Method action.

For example,

- **31512000** | Shaving of benign lesion with chemical cauterization (procedure) | has the Direct morphology of Lesion (morphologic abnormality)

**Direct substance**

Direct substance (attribute) describes the Substance or Pharmaceutical/biologic product on which the procedure's method directly acts.

For example,

- **231274008** | Injection of steroid into joint (procedure) | has Direct substance (attribute) of Steroid (substance)

![Figure 1: Stated view of 231274008 | Injection of steroid into joint (procedure)| with Direct substance (attribute) of Steroid (substance)](image-url)
**Pharmaceutical / biologic product**

Pharmaceutical / biologic product (product) and its descendants are not used as values for Direct Substance (attribute) in the International Release.

Has focus

Has focus (attribute) specifies the Clinical finding or Procedure which is the focus of a procedure. This attribute is self-grouped.

For example,

- 385941006 [Wound care assessment (procedure)] has focus of Wound care (regime/therapy)

Has intent

Has intent (attribute) is used to define procedures that may be done for either a diagnostic or a therapeutic reason. The attribute should be grouped with the attributes that represent the procedure with that intent.

For example,

- 108249004 [Audiologic AND/OR audiometric test including vestibular function (procedure)] is inherently diagnostic, so it would not be modeled with a Has intent (attribute) of Diagnostic intent (qualifier value).
- 274432006 [Therapeutic aspiration of ovary (procedure)] and 274389009 [Diagnostic aspiration of ovary (procedure)] are both modeled with the Has intent (attribute), as the aspiration of ovary procedure can be either therapeutic or diagnostic.

Indirect device

Indirect device (attribute) represents action on something that is located in or on a device, but is not directly on the device itself. This attribute is infrequently needed. When modeling, carefully consider its use.

For example,

- 232762008 [Excision of vegetations from implanted mitral valve (procedure)] has Indirect device of Mitral valve prosthesis device (physical object).

In this example, the vegetation is being excised. The mitral valve prosthesis device is where the vegetation is located, but the mitral valve prosthesis, itself, is not excised. Thus, the mitral valve prosthesis device is the Indirect device.

Indirect morphology

Indirect morphology (attribute) describes the morphology that is acted upon, but is not the direct object of the Method action. This means the procedure acts directly on something else, e.g. a device, substance, or anatomical structure.

For example,

- 404205006 [Removal of mesh from wound (procedure)] has Indirect morphology of Wound (morphologic abnormality)

Method

Method (attribute) represents the action being performed to accomplish the procedure. It does not include: the surgical approach, e.g. translumbar; equipment, e.g. sutures; or physical force, e.g. laser energy (see Surgical Procedures Defining Attribute page).

No relationship group can contain more than one Method (attribute). If a procedure has more than one Method, each serves as the anchor of a separate relationship group that will contain any defining relationships that represent a direct object (and, where relevant, indirect object) of the Method’s action. This is true even if the different Methods each act on the same direct object. Each relationship group can be thought of as representing a component of the procedure that involves a particular action.

For example,

- 10255001 [Incision of ureter (procedure)] has Method (attribute) of Incision - action (qualifier value)
Procedures with a Method (attribute) can be described using an action verb that corresponds to the method. The direct object/s of the action verb should be represented using one or more of the four direct-object attributes, depending on whether the direct object on which the method acts is an:

- Anatomical structure: Procedure site - Direct
- Morphologic abnormality: Direct morphology
- Device: Direct Device
- Substance: Direct Substance

If the anatomical structure, device, or substance of the direct object is indeterminate, do not use the direct-object attributes.

When modeling procedures where the Method is Removal - action (qualifier value) or one of its subtypes, e.g. Excision, Surgical biopsy, etc., for removal of:

- Structures, grafts, and tissue lesions (e.g. cysts, tumors, etc. are considered removal of the site), use Procedure site - Direct.
- Devices, calculi, thrombi, foreign bodies, and other non-tissue entities from the structure, use Procedure site - Indirect.

For example,

- 43748006 |Removal of urinary bladder catheter (procedure)| has a Method (attribute) of Removal - action (qualifier value). Because a device is being removed, use Procedure site - Indirect (attribute) with a value of Bladder and outflow structure (body structure)

**Method attribute**

Attributes should be grouped with the Method attribute to which they apply. In the absence of a Method attribute, related attributes should be grouped together.

Exception,

- Recipient Category (see separate attribute entry)
  - A single procedure concept should not be precoordinated when more than one Recipient Category is involved. Such complex statements should have two or more procedure concepts that are placed into an appropriately structured electronic health application.

**Priority**

Priority (attribute) is used when a procedure concept specifies a priority. This attribute is self-grouped.

For example,

- 708932005 |Emergency hemodialysis (procedure)| has Priority of Emergency (qualifier value)
- 177141003 |Elective cesarean section (procedure)| has Priority of Elective (qualifier value)

260870009 Priority (attribute) is most often used to differentiate elective and emergency subtypes of a procedure that can be performed on either basis. With the exception of Cardiopulmonary resuscitation (procedure), this attribute is normally used only to define concepts whose FSNs specify a priority, not for modeling procedures that imply an emergency priority, such as Heimlich maneuver (procedure) or those that are inherently elective, such as Augmentation mammoplasty (procedure).

**Procedure device**

Procedure device (attribute) is used to model devices associated with a procedure. This attribute is used to define high-level, general concepts that aggregate procedures according to the device involved.

Procedure device subsumes the more specific attributes, Direct Device, Indirect Device, Using Device, and Using Access Device. The more specific attributes should be used instead of Procedure Device, if possible.

For example,

- 276272002 |Catheter procedure (procedure)| has Procedure device of Catheter, device (physical object)
Procedure device

The attribute values in the Procedure Device hierarchy include Device (physical object) and its descendants.

There are a limited number of drug delivery devices in SNOMED CT. These concepts descend from Drug-device combination product (product) which is a descendant of both Device (physical object) and Pharmaceutical / biologic product (product). Although they carry the hierarchy tag of (product), they are acceptable values for attributes in the Procedure Device attribute hierarchy.

Procedure morphology

Procedure morphology (attribute) is used to specify the morphology, or abnormal structure, involved in a procedure. It is used when defining general concepts that subsume direct and indirect morphology. It subsumes the more specific attributes, Direct and Indirect Morphology. These should be used, if possible.

Morphologically abnormal structures

Hematoma, calculus, foreign body, blood clot, embolus, and some other morphologies are not strictly body structures. But, they are included in the body structure hierarchy under morphologically abnormal structure and are valid values for the Procedure Morphology attributes.

Procedure site

Procedure site (attribute) describes the body site acted on or affected by a procedure. The Procedure site (attribute) is

- used to model the site for high-level grouping-type procedure concepts
- most often used for concepts that do not require a Method (attribute) and Action (qualifier value) pair
- not required in order for the classifier to work properly

363704007 | Procedure site (attribute) subsumes the more specific attributes, 405813007 | Procedure site - Direct (attribute), which is the site directly acted upon, and 405814001 | Procedure site - Indirect (attribute), which is the site indirectly acted upon. The more specific attributes should be used if possible (see separate entries for Procedure site - Direct and Procedure site - Indirect).

For example,

- 118839001 | Procedure on colon (procedure) has Procedure site of Colon structure (body structure)

When modeling procedures where the Method is Removal - action (qualifier value) or one of its subtypes, e.g. Excision, Surgical biopsy, etc., for removal of:

- Structures, grafts, and tissue lesions (e.g. cysts, neoplasms, abscesses, wounds, warts, aneurysms, herniations, oral clefts, etc.) are considered removal of the site, use Procedure site - Direct.
- Devices, calculi, thrombi, foreign bodies, and other non-tissue entities from the structure, use Procedure site - Indirect.

Procedure site

Procedures are not necessarily categorized by site.

Use of Structure of <anatomical structure> vs. Entire <anatomical structure> as value of the Procedure site attributes

Structure of <anatomical structure> rather than Entire <anatomical structure> should be used as the value for procedure site attributes, except where the procedure FSN explicitly specified that the entire structure is the object of the procedure.

For example,

- 23968004 | Excision of colon (procedure) has 405813007 | Procedure site - Direct (attribute) of 71854001 | Colon structure (body structure)
• 26390003 | Total colectomy (procedure) | has 405813007 | Procedure site - Direct (attribute) | of 302508007 | Entire colon (body structure)

Procedure site - direct
Procedure site - Direct (attribute) is used when the action of the procedure is directly aimed at anatomical or acquired body structure or site, rather than something else located there (e.g. a device), i.e. when the 260686004 | Method (attribute) | is 129303008 | Removal - action (qualifier value) | or one of its subtypes (Excision, Surgical biopsy, or etc.).

For example,

• 54321008 | Cardiac flow imaging (procedure) | has 405813007 | Procedure site - Direct (attribute) | of Coronary artery structure (body structure)

Tissue lesions (cysts, neoplasms, abscesses, wounds, warts, aneurysms, herniations, oral clefts, etc.) are considered part of the procedure site and should also use 405813007 | Procedure site - Direct (attribute) |.

For example,

• Repair of rectocele (procedure) has a Procedure site - Direct (attribute) of Rectum structure and a Direct morphology (attribute) of Herniated structure
• Closure of skin wound (procedure) has a Procedure site - Direct (attribute) of Skin structure and a Direct morphology (attribute) of Wound
• Fixation of fracture (procedure) has a Procedure site - Direct (attribute) of Bone structure and a Direct morphology (attribute) of Fracture

Procedure site - indirect
Procedure site - Indirect (attribute) specifies the anatomical location but is not the direct focus of the procedure. The direct object of the action may be a device, a substance, or a morphologic abnormality that is not a part of the tissue structure of the anatomical site in which it is located, such as a calculus, thrombus, or foreign body. Thus, 405814001 | Procedure site - Indirect (attribute) | is typically found in a relationship group with a second, "direct" attribute-value relationship, such as a Direct morphology, Direct substance, or Direct device.

For example,

• 405433000 | Removal of catheter from brachial vein (procedure) | has:
  • Method of Removal - action (qualifier value)
  • Procedure site - Indirect of Structure of brachial vein (body structure)
  • Direct device of Venous catheter (physical object)
Figure 2: Stated view of 405433000 |Removal of catheter from brachial vein (procedure)|

- 371005009 |Removal of calculus of urinary bladder (procedure)| has:
  - Direct morphology of Calculus (morphologic abnormality)
  - Method of Removal - action (qualifier value)
  - Procedure site - Indirect of Urinary bladder structure (body structure)
Figure 2: Stated view of 371005009 |Removal of calculus of urinary bladder (procedure)|

Recipient category
Recipient category (attribute) specifies the type of individual or group upon which the action of the procedure is performed.

For example,

- 105455006 |Donor for medical or surgical procedure (person)| has Recipient Category (attribute) of Donor if the subject of the record is the Blood product donor (person).
  
  This can be used in blood banking procedures to differentiate the donor vs the recipient of blood products.

⚠ **Recipient category**

It is not used for a procedure where the subject of the procedure is someone other than the subject of record.

Revision status
Revision status (attribute) refers to another procedure performed on the same site for the same condition. A procedure without a revision status is considered to be performed for the first time. A revision procedure can be modeled with a Revision status (attribute) of Revision - value (qualifier value).

For example,

- 128323000 |Revision of implant (procedure)| has Revision status of Revision - value (qualifier value)

Using access device
Using access device (attribute) specifies the instrument or equipment used to access the site of a procedure.

For example,

- 301761003 |Arthroscopic synovial biopsy (procedure)| has Using access device of Arthroscope, device (physical object)
Using device (attribute) refers to the instrument or equipment utilized to execute an action. It is used when the device is actually used to carry out the action, that is the focus of the procedure. If the device is simply the means to access the site of the procedure, then Using access device is the appropriate attribute.

For example,
- 51064005 | Core needle biopsy of larynx (procedure) has Using device of Core biopsy needle, device (physical object)

Using energy
Using energy (attribute) refers to the energy used to execute an action.

For example,
- 65952009 | Gamma ray therapy (procedure) has Using energy of Gamma radiation (physical force)

Using substance
Using substance (attribute) describes the Substance used to execute the action of a procedure. It is not the substance on which the procedure's method directly acts, the Direct substance.

For example,
- 285754008 | Contrast radiography of esophagus (procedure) has Using substance of Contrast media (substance)

Evaluation Procedure
The following defining attributes are unique in the context of the 386053000 | Evaluation procedure (procedure) | subhierarchy. Many of these attributes (e.g., Component, Scale type) are used to define Observable entity concepts. Evaluation procedures may use the attributes below in addition to those attributes allotted to the 71388002 | Procedure (procedure) | hierarchy (see Procedure Attributes Summary page). All of the attributes for Evaluation procedure concepts should be grouped. However, current modeling does not reflect this guidance, and examples used on this page are inconsistent with current guidance. The future of this hierarchy in relation to the observable entity hierarchy is under review. See 'Observable Entity vs. Evaluation Procedure' at Observable Entity.

Component
Component refers to what is being observed or measured by a procedure.

For example,
- 442165003 | Quantitative measurement of polychlorinated biphenyl in blood specimen using gas chromatography (procedure) has 246093002 | Component (attribute) of 42001007 | Polychlorinated biphenyl (substance)
Has specimen

*Has Specimen* indicates the type of specimen on which a measurement or observation is performed.

For example,

- `442165003` [Quantitative measurement of polychlorinated biphenyl in blood specimen using gas chromatography (procedure)] uses `116686009` [Has specimen (attribute)] of `119297000` [Blood specimen (specimen)]

Measurement method

*Measurement Method* specifies the method by which an evaluation procedure is performed. It provides additional specificity. For measurement procedures, the `260686004` [Method (attribute)] is given the value `129266000` [Measurement - action (qualifier value)]. No concept can be defined with a `370129005` [Measurement method (attribute)] unless it is being used to refine a `260686004` [Method (attribute)] that has a value of `129266000` [Measurement - action (qualifier value)] or one of its subtypes that is also specified in the concept definition. That is, use of `370129005` [Measurement method (attribute)] must be in addition to a `260686004` [Method (attribute)] of `129266000` [Measurement - action (qualifier value)] or one of its subtypes. Also, the `370129005` [Measurement method (attribute)] and its value must be grouped with the `260686004` [Method (attribute)] and its value of the concept or subtype of `129266000` [Measurement - action (qualifier value)].

For example,

- `442165003` [Quantitative measurement of polychlorinated biphenyl in blood specimen using gas chromatography (procedure)] has a `370129005` [Measurement method (attribute)] of `2842000` [Gas chromatography measurement (procedure)]

Property

*Property* specifies the kind of property (quality or characteristic) being measured.

For example,

- `19165008` [Measurement of limb length (procedure)] has a `370130000` [Property (attribute)] of `410668003` [Length property (qualifier value)]
Scale type

Scale Type refers to the scale of the result of an observation of a diagnostic test.

For example,

- 442165003 [Quantitative measurement of polychlorinated biphenyl in blood specimen using gas chromatography (procedure)] has 370132008 [Scale type (attribute)] of 30766002 [Quantitative (qualifier value)]

Time aspect

Time Aspect specifies temporal relationships for a measurement procedure. While this attribute has been approved, guidelines for its implementation await development.

Further clarification

An evaluation procedure may evaluate a property of a component, or a property may be the sole focus of the method. In the latter case, component isn’t included since only the property is being evaluated.

For example of an evaluation procedure evaluating a property of a component,

- 443834000 [Quantitative measurement of mass concentration of bismuth in urine specimen (procedure)] has 370130000 [Property (attribute)] of 118539007 [Mass concentration (property) (qualifier value)], and 246093002 [Component (attribute)] of 23172004 [Bismuth (substance)]
For example where property may be the sole focus of the method,

- 78888000 | Osmolality measurement, urine (procedure) |
Surgical Procedure

The following defining attribute is unique to Surgical procedures. Surgical procedures may also use the attributes in the Procedure Attributes Summary table from the HRCM (see also Procedure Defining Attributes page).

Surgical approach

Surgical Approach specifies the directional, relational, or spatial access to the site of a surgical procedure. The range for Surgical Approach is descendants of 103379005 | Procedural approach (qualifier value)|.

- 172883004 | Intranasal ethmoidectomy (procedure)| has Surgical approach, Intranasal approach (qualifier value)

Administration of Substance via Specific Route Procedure

In addition to attributes applicable to general procedures, the subhierarchy of |Administration of substance via specific route (procedure)| also includes the |Route of administration (attribute)|.

Route of administration

Route of administration represents the route by which a procedure introduces a substance into the body. The domain for this attribute is descendants of 433590000 | Administration of substance via specific route (procedure)|. The range involves subtypes of 284009009 | Route of administration value (qualifier value)|. When using this attribute, an additional attribute of |Procedure site - indirect| should be modeled and grouped with the |Route of administration (attribute)|.

For example,

- 410572008 | Intravitreal steroid injection (procedure)| has the | Route of administration (attribute) | of Intravitreal route (qualifier value)
While the values for the |Procedure site - Indirect| and |Route of administration| attributes may be similar and seem redundant, their presence is necessary for consistent subsumption.

<table>
<thead>
<tr>
<th>Route of administration (qualifier value)</th>
<th>Body structure value of Procedure site - Indirect</th>
</tr>
</thead>
<tbody>
<tr>
<td>Intravenous route</td>
<td>Venous structure</td>
</tr>
<tr>
<td>Oral route</td>
<td>Mouth region structure</td>
</tr>
</tbody>
</table>

Table 1: Examples of complementary values of Route of administration and Procedure site - Indirect

Procedure Naming Conventions

General rules

The naming pattern for procedures is highly dependent on the attributes used to describe the procedure. However, some general rules apply.

The FSN for a procedure should name the action (the method) of the procedure first, and then the object that the action directly acts upon.

For example,

- 261531000 |Excision of rib (procedure)|

Include the approach where more than one option exists. See the various approaches as subtypes of 103379005 |Procedural approach (qualifier value)|.

For example,

- 386792000 Transurethral resection of neoplasm of bladder (procedure) vs 287726000 Open resection of neoplasm of bladder (procedure)
Because a resection of a neoplasm of the bladder can be performed by transurethral and open approaches, concepts are separately identified with each approach.

The preferred term should match the FSN. Synonyms using the common clinical names of procedures are acceptable.

A common naming pattern for concepts in the procedure hierarchy is:

FSN: <Method (Action)> of <Anatomical or acquired body structure> (procedure)
PT:  <Method (Action)> of <Anatomical or acquired body structure>
SYN: [plasty/stomy/ectomy/otomy]

For example,  
• 82035006 | Resection of polyp (procedure) | has a synonym of polypectomy

Further refinements can be affected by the various attributes and their values as described in the sections below.

Anatomical site
An anatomical site is the direct object of the action. The name of the site should follow the name of the action.

For example,  
• 175253007 | Repair of pulmonary artery (procedure) |
  The action is repair and the site is pulmonary artery. The action is listed first in the description, followed by the site.

Procedure descriptions should follow the naming guidelines for the body structure hierarchy. Concepts describing limbs are frequently found in SNOMED CT, and the use of upper/lower limb in the FSN with synonyms of upper/lower extremity, arm/leg should be followed.

For example,  
• 179987000 | Replantation of upper limb (procedure) |
  The description of upper limb is used in the FSN while the synonyms refer to arm and upper extremity.

Device
A device is the direct object of the action. The word(s) naming the device should follow the word(s) naming the action. If there is a site that is not the direct object of the action, the word(s) naming it should come after the word(s) naming the device.

For example,  
• 392247006 | Insertion of catheter into artery (procedure) | The action is insertion, the direct object is catheter, and the indirect site is artery.
Substance

A **substance** is the direct object of the action. The word(s) that name the substance should follow the words that name the action. If there is a site that is not the direct object of the action, the word(s) naming it should follow the word(s) naming the substance.

For example,

- **427258004 Injection of hormone into subcutaneous tissue (procedure)** The action is *injection*, the direct object is *hormone*, and the indirect site is *subcutaneous tissue*.

Morphologic abnormality

A **morphologic abnormality** is the direct object of the action. The morphology term should follow the action term. If there is a site, it should follow the morphology term.

For example,

- **41180005 Excision of cyst of breast (procedure)** The action is *excision*, the direct object is the morphologic abnormality *cyst*, and the site is *breast*.
- **175376008 Operation on aneurysm of carotid artery (procedure)** The action is *operation*, the direct object is the morphologic abnormality *aneurysm*, and the site is *carotid artery*.

Past tense verbs and sentence types

A procedure concept should be a noun phrase that names the procedure. It should not contain information that it was done, or is to be ordered, carried out, or planned.

- Past tense verbal phrases should not be used to name procedures, since *past tense* invokes a temporal context, i.e. the procedure was done in the past. Any existing concepts with past tense verbs should be moved to the Situation with explicit context hierarchy.
- Sentence function types, i.e. imperative, declarative, interrogative, or exclamatory, are disallowed in procedure concepts.

Acceptable example,

- **11227005 Excision of ganglion of tendon sheath of hand (procedure)** This is an acceptable FSN expressed with a noun phrase.

Unacceptable example,

- *Hand tendon ganglion excised* indicates the procedure was done, as a past tense declarative statement. This should be in the Situation with explicit context hierarchy, not the Procedure hierarchy.

Complexity

Complexity can mean either the amount of effort required, or it can be based on realm-specific definitions (e.g. simple arthrodesis, simple repair, complex repair, etc.). Procedure concepts with modifiers representing complexity are not allowed in the International Release.

**Exception**

Procedures that use the terms *simple* or *complex* are allowed if defined with reproducible meanings, based on what is done to or for the patient, rather than how much effort is expended.

For example,

- **172043006 Simple mastectomy (procedure)** The concept is reproducibly defined as the removal of all breast tissue without removal of axillary contents. This is differentiated from modified radical, radical, skin-sparing, and subcutaneous variants of mastectomy.
Procedures by count

Counts of the number of procedures
Many procedure classifications focus on resources required to complete; this may be for reimbursement or tracking purposes (e.g. placement of one stent versus placement of two stents). This information should be part of patient documentation and is not allowed in the International Release.

Order of procedures
The order of procedures, e.g. primary, first, second, etc. should be excluded.

Bilateral Procedure Naming Conventions

The acceptable naming pattern for procedures with lateralizable body parts:

- FSN: <Method> of bilateral <anatomical or acquired body structure> (procedure)
- PT:  <Method> of bilateral <anatomical or acquired body structure>
- SYN: <Method> of both <anatomical or acquired body structure>

For example,

895470004 | Amputation of bilateral upper limbs (procedure) |

FSN: Amputation of bilateral upper limbs (procedure)  
PT:  Amputation of bilateral upper limbs  
SYN: Amputation of both upper limbs
Chemotherapy Regime Therapy Naming Conventions

Chemotherapy regimens, which are internationally recognized and implemented, are acceptable content and may be added to the (regime/therapy) hierarchy as subtypes of 716872004 |Antineoplastic chemotherapy regimen (regime/therapy)|.

Trade names, which are indicated by an acronym, e.g. ABVD chemotherapy regimen, where A represents trade name Adriamycin®, should not be spelled out but may be referenced in the acronym used to describe the regimen.

Generic drug names are not capitalized; they are lower case. When creating descriptions for generic drug names, the substance described should match the description from the Substance hierarchy.

Examples,

**ABVD chemotherapy regimen.** A represents the trade name Adriamycin® (INN = doxorubicin):
- FSN: Doxorubicin, bleomycin, vinblastine and dacarbazine chemotherapy regimen (regime/therapy)
- PT: ABVD chemotherapy regimen
- Synonym: Doxorubicin, bleomycin, vinblastine and dacarbazine chemotherapy regimen
- Synonym: ABVD chemotherapy protocol

**R-CHOP chemotherapy regimen.** H represents the non-INN generic name hydroxydaunomycin (INN = doxorubicin) and O represents the trade name Oncovin® (INN = vincristine):
- FSN: Rituximab, cyclophosphamide, doxorubicin, vincristine and prednisone chemotherapy regimen (regime/therapy)
- PT: R-CHOP chemotherapy regimen
- Synonym: Rituximab, cyclophosphamide, doxorubicin, vincristine and prednisone chemotherapy regimen
- Synonym: R-CHOP chemotherapy protocol

Other synonyms may be added if requested, e.g. *Left and right X; Bilateral X*, etc.
Clinical Imaging Procedure Naming Conventions

Almost all imaging procedures can be unambiguously expressed in a number of ways. There is a balance between flexibility in language and efficiency in terminology maintenance. Consequently, all variants for imaging modalities are not routinely included in SNOMED CT. Submissions for additional descriptions must be justified explicitly.

At a minimum, procedures are ordinarily expressed with the modality and body site. Existing content may have inconsistencies, but new content should follow the naming conventions that follow.

- **The use of near synonyms is acceptable for clinical imaging procedures:**

  - For example,
    - 79516005 Renal arteriography (procedure) has the synonym renal angiography
    - 726077005 Computed tomography arteriography of bronchial artery (procedure) has the synonym CT angiography of bronchial artery
    - 709552006 Computed tomography angiography of iliac artery (procedure) has the synonym CT angiogram of iliac artery

- **Under revision**

  There is inconsistency with naming Radiology of X vs X-ray of X and modeling of X-ray concepts. Preliminary analysis has been completed and a new approach recommended. Remodeling is pending.

- **X-ray**

  **Approach 1:** Radiography of X
  - FSN: Radiography of X (procedure)
  - PT: Radiography of X
  - For example,
    - 49345004 Radiography of hand (procedure)

  **Approach 2:** X-ray of X
  - FSN: X-ray of X (procedure)
  - PT: X-ray of X
  - For example,
    - 426581005 X-ray of both feet (procedure)

- **Diagnostic radiography**

  363680008 | Radiographic imaging procedure (procedure) | is at the top-level of the hierarchy of imaging procedures utilizing X-rays. The phrase diagnostic radiography is allowed as an FSN of subtypes of radiographic imaging procedure.

  - For example,
    - 66596009 Diagnostic radiography for foreign body detection and localization (procedure)

- **Inactivated concept**

  Diagnostic radiologic examination (procedure) had a synonym of X-ray. It may have been interpreted more narrowly, because of the potential for a narrower interpretation of radiologic vs. radiographic and diagnostic.

- **Modeling: New content requests**

  An X-ray concept may have the action, 312254007 Plain X-ray imaging - action (qualifier value) or the broader supertype action, 278110001 Radiographic imaging - action (qualifier value). A submitter should clearly identify which of the actions is appropriate.

Ultrasonography
Ultrasonography
   • FSN: Ultrasonography of X (procedure)
   • PT: Ultrasonography of X
   • SYN: Ultrasound scan of X
   • SYN: Ultrasound of X
   For example,
     • 709590000 [Ultrasonography of perineum (procedure)]
       • FSN: Ultrasonography of perineum (procedure)
       • PT: Ultrasonography of perineum
       • SYN: Ultrasound scan of perineum
       • SYN: Ultrasound of perineum

Doppler ultrasonography
   • FSN: Doppler ultrasonography of X (procedure)
   • PT: Doppler ultrasonography of X
   • SYN: Doppler ultrasound scan of X
   • SYN: Doppler ultrasound of X
   For example,
     • 710306004 [Doppler ultrasonography of venous structure (procedure)]
       • FSN: Doppler ultrasonography of venous structure (procedure)
       • PT: Doppler ultrasonography of vein
       • SYN: Doppler ultrasound scan of vein
       • SYN: Doppler ultrasound of vein

Obstetric ultrasonography
An obstetric ultrasound may require a complex description. However, the same rules apply, as follows:
   • FSN: Obstetric ultrasonography of X (procedure)
   • PT: Obstetric ultrasonography of X
   • SYN: Obstetric ultrasound scan of X
   • SYN: Obstetric ultrasound of X
   For example,
     • 169670003 [Antenatal ultrasound scan at 17-22 weeks (procedure)]
       • FSN: Antenatal ultrasound scan at 17-22 weeks (procedure)
       • PT: Antenatal ultrasound scan at 17-22 weeks

Computed Tomography - CT

<i>Exception for abbreviation</i>

CT is an exception to the rule that all abbreviations should have their expanded form in parentheses in descriptions.
Axial & Scan

Legacy issues: Existing computerized tomography concepts will be renamed consistently when the Quality Initiative undertakes improvement of the procedure hierarchy.

Axial

Requests for new descriptions with computerized axial tomography (CAT) are not acceptable. The axial part of the phrase is no longer accurate, because there are other techniques that also create images on multiple planes or axes.

Scan

Computed tomography descriptions do not routinely include computed tomography scan of X.

The word scan is not systematically added in new descriptions and should not be included in preferred terms. However, specific requests to add descriptions with the word scan are not denied.

Computed tomography angiography scan of X is considered obsolescent and should not be added as a new description.

With Contrast

CT angiography uses contrast in all cases of the procedure; this must be stipulated in the descriptions.

Computed tomography angiography with contrast

This naming pattern is used when an anatomical location is specified, but the blood vessel is not explicit.

- FSN: Computed tomography angiography of X with contrast (procedure)
- PT: CT angiography of X with contrast
- SYN: Computed tomography angiography of X with contrast

For example,

- 582101000119108 |Computed tomography angiography of head with contrast (procedure)|
  - FSN: Computed tomography angiography of head with contrast (procedure)
  - PT: CT angiography of head with contrast
  - SYN: Computed tomography angiography of head with contrast

Computer tomography arteriography with contrast

- FSN: Computed tomography arteriography of [artery] with contrast (procedure)
- PT: CT arteriography of [artery] with contrast
- SYN: Computed tomography arteriography of [artery] with contrast
- SYN: CT arteriogram of [body structure] with contrast
- SYN: CT arteriography of [body structure] with contrast

For example,

- 726077005 Computed tomography angiography of bronchial artery with contrast (procedure)
  - FSN: Computed tomography angiography of bronchial artery with contrast (procedure)
  - PT: CT angiography of bronchial artery with contrast
• SYN: CT arteriography of bronchial artery with contrast
• SYN: Computed tomography arteriography of bronchial artery with contrast

Computed tomography venography with contrast
• FSN: Computed tomography venography of X with contrast (procedure)
• PT: CT venography of X with contrast
• SYN: CT venogram of X with contrast
• SYN: Computed tomography venography of X with contrast
For example,
  • 432842007 |Computed tomography venography of intracranial vein with contrast (procedure)|
    • FSN: Computed tomography venography of intracranial vein with contrast (procedure)
    • PT: CT venography of intracranial vein with contrast
    • SYN: Computed tomography venography of intracranial vein with contrast

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Venography

Venography may simply be a timing phase of angiography. It is agreed that venography may be a useful term in an FSN, i.e. there may be a meaningful technique difference between simple angiography and purposeful venography.

Computed tomography arthrography
• FSN: Computed tomography arthrography of X (procedure)
• PT: CT arthrography of X
• SYN: CT arthrogram of X
• SYN: Computed tomography arthrography of X
For example,
  • 418940000 |Computed tomography arthrography of intratarsal joint (procedure)|
    • FSN: Computed tomography arthrography of intratarsal joint (procedure)
    • PT: CT arthrogram of intratarsal joint
    • SYN: CT arthrography of intratarsal joint

Magnetic Resonance Imaging - MRI

Magnetic resonance imaging
Descriptions:
• FSN: Magnetic resonance imaging of X (procedure)
• PT: MRI of X
• SYN: Magnetic resonance imaging of X
For example,
  • 6007000 |Magnetic resonance imaging of chest (procedure)|
    • FSN: Magnetic resonance imaging of chest (procedure)
    • PT: MRI of chest
    • SYN: Magnetic resonance imaging of chest

Magnetic resonance angiography
Descriptions:
• FSN: Magnetic resonance angiography of X (procedure)
• PT: Magnetic resonance angiography of X
• SYN: Magnetic resonance angiogram of X
• SYN: MR angiography of X

---
For example,

- 432103005 | Magnetic resonance angiography of carotid artery (procedure) |
  - FSN: Magnetic resonance angiography of carotid artery (procedure)
  - PT: Magnetic resonance angiography of carotid artery
  - SYN: Magnetic resonance angiogram of carotid artery
  - SYN: MR angiography of carotid artery

Magnetic resonance venography

Descriptions:

- FSN: Magnetic resonance venography of X (procedure)
- PT: Magnetic resonance venography of X
- SYN: Magnetic resonance venogram of X
- SYN: MR venography of X

For example,

- 21101000087100 | Magnetic resonance venography of limb (procedure) |
  - FSN: Magnetic resonance venography of limb (procedure)
  - PT: Magnetic resonance venography of extremity
  - SYN: Magnetic resonance venography of limb

Magnetic resonance arthrography

Descriptions:

- FSN: Magnetic resonance arthrography of X (procedure)
- PT: Magnetic resonance arthrography of X
- SYN: Magnetic resonance arthrogram of X
- SYN: MR arthrography of X

For example,

- 19741000087109 | Magnetic resonance arthrography of right knee (procedure) |
  - FSN: Magnetic resonance arthrography of right knee (procedure)
  - PT: Magnetic resonance arthrography of right knee
  - SYN: MR arthrography of right knee

Magnetic resonance spectroscopy

Descriptions:

- FSN: Magnetic resonance spectroscopy of X (procedure)
- PT: Magnetic resonance spectroscopy of X
- SYN: MR spectroscopy of X

For example,

- 1137352004 | Magnetic resonance spectroscopy of liver (procedure) |
  - FSN: Magnetic resonance spectroscopy of liver (procedure)
  - PT: Magnetic resonance spectroscopy of liver
  - SYN: MR spectroscopy of liver

Contrast for Imaging

It is essential to express when contrast is part of a procedure and that descriptions are constructed consistently.

For example,

- 702501008 | Computed tomography of knee with contrast (procedure) |
  - FSN: Computed tomography of knee with contrast (procedure)
  - PT: CT of knee with contrast
  - SYN: Computed tomography of knee with contrast

The descriptions for all procedure concepts that use contrast must explicitly state ‘with contrast.’ All procedures that use contrast, including 420040002 | Fluoroscopic angiography with contrast (procedure) | and all subtypes, must
include the attribute 424361007 |Using substance (attribute)| and value 385420005 |Contrast media (substance)| in the role group.

It is unnecessary to add the word media to contrast.

It is agreed that the linking word to associate the contrast use with the procedure is with not for, or, etc.

### Contrast

There is a suggestion that additional concept detail is required when it is necessary to know the more precise nature of contrast (e.g. iodinated with various osmolalities, barium, or gas).

### Imaging without contrast

Although considered a negation, this term is used in clinical records. Without contrast imaging procedures are acceptable.

For example,

- 566341000119106 |Computed tomography of ankle without contrast (procedure)|
  - FSN: Computed tomography of ankle without contrast (procedure)
  - PT: CT ankle without contrast
  - SYN: Computed tomography of ankle without contrast

### Without contrast

There is a case for explicitly adding a concept qualification when naming procedures that are explicitly performed without contrast.

In the UK and Australia, it was reported that there are no procedures that specify without contrast pre-coordinated in the national subset. With this information, implementation guidance may be provided.

### Unacceptable concept qualification

With and without and With or without imaging concepts are not acceptable due to ambiguity. Two concepts should be used to express these separately.

### Imaging Guided Procedures

There are numerous procedures where the imaging component is considered a supplemental or secondary technique to help accomplish the primary goal. The pattern is:

#### Procedure using guidance

- FSN: Y (procedure) using (DI Modality guidance) (procedure)
- PT: (DI Modality) guided Y (procedure)
- SYN: Y (procedure) using (DI Modality guidance)

For example,

- 407971000119109 |Percutaneous needle biopsy of liver using computed tomography guidance (procedure)|
  - FSN: Percutaneous needle biopsy of liver using computed tomography guidance (procedure)
  - PT: CT guided biopsy of liver
  - SYN: Percutaneous needle biopsy of liver using computed tomography guidance

#### Computed tomography guided procedure

- FSN: Y using computed tomography guidance (procedure)
- PT: CT guided Y
- SYN: Y using computed tomography guidance

For example,

- 431864000 |Injection using computed tomography guidance (procedure)|
  - FSN: Injection using computed tomography guidance (procedure)
  - PT: CT guided injection
• SYN: Injection using computed tomography guidance

Fluoroscopy guided procedure
• FSN: Y using fluoroscopic guidance (procedure)
• PT: Fluoroscopy guided Y
• SYN: Y using fluoroscopic guidance

For example,
• 773291002 | Biopsy of abdomen using fluoroscopic guidance (procedure) |
  • FSN: Biopsy of abdomen using fluoroscopic guidance (procedure)
  • PT: Fluoroscopy guided biopsy of abdomen
  • SYN: Biopsy of abdomen using fluoroscopic guidance

**Fluoroscopic guidance**
The term fluoroscopic Y is interpreted as Y using fluoroscopic guidance (procedure). Procedures such as 432540009 | Biopsy of wrist using fluoroscopic guidance (procedure) | are subtypes of Fluoroscopy (procedure).

(See also Fluoroscopy and Fluoroscopic Imaging page)

Magnetic resonance imaging guided procedure
• FSN: Y using magnetic resonance imaging guidance (procedure)
• PT: MRI guided Y (procedure)
• SYN: Y using magnetic resonance imaging guidance

For example,
• 433008009 | Core needle biopsy of breast using magnetic resonance imaging guidance (procedure) |
  • FSN: Core needle biopsy of breast using magnetic resonance imaging guidance (procedure)
  • PT: MRI guided core needle biopsy of breast
  • SYN: Core needle biopsy of breast using magnetic resonance imaging guidance

Ultrasonography guided procedure
• FSN: Y using ultrasonographic guidance (procedure)
• PT: Ultrasonography guided Y
• SYN: Y using ultrasonographic guidance

For example,
• 710790002 | Puncture and aspiration using ultrasonographic guidance (procedure) |
  • FSN: Puncture and aspiration using ultrasonographic guidance (procedure)
  • PT: Ultrasonography guided puncture and aspiration
  • SYN: Puncture and aspiration using ultrasonographic guidance

X-ray guided procedure
• FSN: Y using X-ray guidance (procedure)
• PT: X-ray guided Y
• SYN: Y using X-ray guidance

For example,
• 718674009 | Injection of steroid using X-ray guidance (procedure) |
  • FSN: Injection of steroid using X-ray guidance (procedure)
  • PT: X-ray guided steroid injection
  • SYN: Injection of steroid using X-ray guidance

Fluoroscopy and Fluoroscopic Imaging

Simple fluoroscopy
Simple fluoroscopy is real time imaging (usually on TV monitors/image intensifiers) of a body part or system. Only rarely is it an imaging process alone (without some interventional procedure). Fluoroscopy is most often used to guide or direct a primary procedure/purpose. The usual convention in clinical practice is to ignore the fluoroscopic
element and refer to a procedure entirely by the primary component, e.g. angiography. However, this is unacceptable in SNOMED CT, where the imaging component must be explicitly described.

- FSN: Fluoroscopy of X (procedure)
- PT: Fluoroscopy of X
- SYN: Fluoroscopy - X

For example,

- 169005008 | Fluoroscopy of esophagus (procedure) |
  - FSN: Fluoroscopy of esophagus (procedure)
  - PT: Fluoroscopy of esophagus
  - SYN: Fluoroscopy - esophagus

Fluoroscopic guidance

Fluoroscopic X is interpreted as X using fluoroscopic guidance (procedure). Such procedures are subtypes of Fluoroscopy (procedure). See also Imaging modeling guidance.

- FSN: X using fluoroscopic guidance (procedure)
- PT: Fluoroscopy guided X
- SYN: X using fluoroscopic guidance

For example,

- 710293001 | Colonoscopy using fluoroscopic guidance (procedure) |
  - FSN: Colonoscopy using fluoroscopic guidance (procedure)
  - PT: Fluoroscopy guided colonoscopy
  - SYN: Colonoscopy using fluoroscopic guidance

For example,

- Angioplasty using fluoroscopic guidance with contrast (procedure)
  - FSN: Angioplasty using fluoroscopic guidance with contrast (procedure)
  - PT: Fluoroscopy guided angioplasty with contrast
  - SYN: Angioplasty using fluoroscopic guidance with contrast
  - SYN: Fluoroscopic angioplasty with contrast

Fluoroscopic angiography with contrast

Angiography and angiogram, as terms on their own, refer to visualization of a blood vessel or vascular structure not specified as an artery or a vein and could refer to either or both. So for clarity, a blood vessel site (X) and the phrase 'with contrast' must be included in the concept description.

Fluoroscopic angiography always uses contrast. 'With contrast' must be explicitly stated in all descriptions.

Fluoroscopic angiography when vessel is not stated

- FSN: Fluoroscopic angiography of X with contrast (procedure)
- PT: Fluoroscopic angiography of X with contrast
- SYN: Fluoroscopic angiogram of X with contrast

Fluoroscopic angiography of artery

- FSN: Fluoroscopic angiography of X artery with contrast (procedure)
- PT: Fluoroscopic angiography of X artery with contrast
- SYN: Fluoroscopic arteriography of X with contrast
- SYN: Fluoroscopic arteriogram of X with contrast

For example,

- FSN: Fluoroscopic angiography of right cervical vertebral artery with contrast (procedure)
- PT: Fluoroscopic angiography of right cervical vertebral artery with contrast
- SYN: Fluoroscopic arteriography of right cervical vertebral artery with contrast
• SYN: Fluoroscopic arteriogram of right cervical vertebral artery with contrast

Fluoroscopic venography
• FSN: Fluoroscopic venography of X with contrast (procedure)
• PT: Fluoroscopic venography of X with contrast
• SYN: Fluoroscopic venogram of X with contrast

For example,
• FSN: Fluoroscopic venography of right extremity with contrast (procedure)
• PT: Fluoroscopic venography of right extremity with contrast
• SYN: Fluoroscopic venogram of right extremity with contrast

Fluoroscopic arthrography
• FSN: Fluoroscopic arthrography of X (procedure)
• PT: Fluoroscopic arthrography of X
• SYN: Fluoroscopic arthrogram of X

For example,
• 723776000 Fluoroscopic arthrography of right sacroiliac joint (procedure)
  • FSN: Fluoroscopic arthrography of right sacroiliac joint (procedure)
  • PT: Fluoroscopic arthrography of right sacroiliac joint
  • SYN: Fluoroscopic arthrogram of right sacroiliac joint

Dual energy X-ray photon absorptiometry
• FSN: Dual energy X-ray photon absorptiometry of X (procedure)
• PT: Dual energy X-ray photon absorptiometry of X
• SYN: DXA of X
• SYN: DEXA of X

For example,
• 722297000 Dual energy X-ray photon absorptiometry of vertebral column (procedure)
  • FSN: Dual energy X-ray photon absorptiometry of vertebral column (procedure)
  • PT: Dual energy X-ray photon absorptiometry of vertebral column
  • SYN: DXA of vertebral column
  • SYN: DEXA of vertebral column

Positron Emission Tomography - PET

Positron emission tomography (procedure)
• FSN: Positron emission tomography of X (procedure)
• PT: PET of X
• SYN: Positron emission tomography of X

For example,
• 702577007 Positron emission tomography of whole body (procedure)
  • FSN: Positron emission tomography of whole body (procedure)
  • PT: PET of whole body
  • SYN: Positron emission tomography of whole body

Single Photon Emission Computed Tomography - SPECT

Single photon emission computed tomography (procedure)
• FSN: Single photon emission computed tomography of X (procedure)
• PT: Single photon emission computed tomography of X
• SYN: SPECT of X

For example,
• 709549003 Single photon emission computed tomography of heart (procedure)
  • FSN: Single photon emission computed tomography of heart (procedure)
  • PT: Single photon emission computed tomography of heart
Syn: SPECT of heart

Multi-modality Imaging: PET, CT and SPECT, CT

There are very few imaging procedures which are truly multi-modality procedures. Two procedures are usually conducted in parallel, rather than as one. Positron emission tomography with computed tomography (PET/CT) and Single photon emission computed tomography with computed tomography (SPECT/CT), however, are produced by one piece of equipment, possibly by a single technician, but with multiple imaging energies.

Positron emission tomography with computed tomography

- FSN: Positron emission tomography with computed tomography of X (procedure)
- PT: PET CT of X
- SYN: Positron emission tomography with computed tomography of X

For example,

- 16554361000119106 | Positron emission tomography with computed tomography of brain (procedure)
  - FSN: Positron emission tomography with computed tomography of brain (procedure)
  - PT: PET CT of brain
  - SYN: Positron emission tomography with computed tomography of brain

Single photon emission computed tomography with computed tomography

- FSN: Single photon emission computed tomography with computed tomography of X (procedure)
- PT: Single photon emission computed tomography with computed tomography of X
- SYN: SPECT CT of X

For example,

- 16534151000119105 | Single photon emission computed tomography with computed tomography of liver (procedure)
  - FSN: Single photon emission computed tomography with computed tomography of liver (procedure)
  - PT: Single photon emission computed tomography with computed tomography of liver
  - SYN: SPECT CT of liver

Nuclear Medicine - Radionuclide Imaging

Nuclear medicine imaging uses radionuclides or radioisotopes.

Radionuclide scan

- FSN: Radionuclide scan of X (procedure)
- PT: Radionuclide scan of X
- SYN: Radioisotope scan of X

For example,

- 710313004 | Radionuclide scan of peritoneal cavity (procedure)
  - FSN: Radionuclide scan of peritoneal cavity (procedure)
  - PT: Radionuclide scan of peritoneal cavity
  - SYN: Radioisotope scan of peritoneal cavity

Radionuclide scan using isotopes (with other agents)

- FSN: Radionuclide scan of X using Y (procedure)
- PT: Radionuclide scan of X using Y
- SYN: Radioisotope scan of X using Y

For example,

- 710312009 | Radionuclide scan of perfusion of liver using technetium Tc^99m^ aggregated albumin (procedure)
  - FSN: Radionuclide scan of perfusion of liver using technetium Tc^99m^ aggregated albumin (procedure)
  - PT: Radionuclide scan of perfusion of liver using technetium Tc^99m^ aggregated albumin
  - SYN: Radioisotope scan of perfusion of liver using technetium Tc^99m^ aggregated albumin
Diagnostic Imaging for Multiple Body Sites

Adjacent structures

Concepts which describe adjacent structures, imaged in one procedure, are acceptable.

For example,

- 432672003 |Magnetic resonance imaging of pelvis and hip (procedure)|

**Unacceptable**

Multiple procedures or a combination of different procedures in one concept are unacceptable.

Unacceptable examples,

- Computed tomography angiography of aorta, abdomen, pelvis and lower limb
- Ultrasonography of abdomen and ultrasonography of pelvis with transrectal ultrasonography
- Ultrasonography of pelvis and obstetric ultrasonography with transvaginal ultrasonography
- Ultrasonography of knee and Doppler ultrasonography of vein of lower limb

Imaging Adjustments for View, Projection, or Technique

It may be important, from both clinical and administrative perspectives, to capture variations or modifications of imaging technique. The variations may impact correct acquisition and interpretation of images.

Examples of modifications include:

- Axial (qualifier value)
- Skyline projection (qualifier value)
- Decubitus (qualifier value)

**Post-coordination**

Though these examples are qualifying values in SNOMED CT, they are not allowable for post-coordination of diagnostic imaging procedures.

Measurement Procedure Naming Conventions

SNOMED International is no longer accepting new concepts to the 122869004 |Measurement procedure (procedure)| hierarchy. New requests for content in this area will be created in the 363787002 |Observable entity (observable entity)| hierarchy. Legacy content will be relocated from the Measurement (procedure) hierarchy to the Observable entity hierarchy in the future. Please see Observable Entity and Observable Entity Naming Conventions pages for more information.

Procedure Modeling

Procedure attribute hierarchies

SNOMED CT has attribute hierarchies for Procedure Site, Procedure Device, and Procedure Morphology. Each has two sub-attributes to represent the direct and indirect objects. Procedure Device also has more specific attributes, Using Device and Using Access Device.
Observable Entity vs. Evaluation Procedure

The observable entity and evaluation procedure hierarchies have some of the same attributes. There is not and should not be a one-to-one correspondence between the two hierarchies.

Concepts will not be duplicated between the observable entity hierarchy and procedure hierarchy, and requests for such will not be added. While some users have indicated they want to use a procedure concept for ordering a test and an observable concept for reporting the result, this is not an acceptable use case.

At this time, SNOMED CT contains some concepts in the procedure hierarchy which logically belong in the observable entity hierarchy. It is noted that these concepts will likely move to the observable entity hierarchy in the future. In addition, if we identify existing duplicate concepts between the two hierarchies, this will also be corrected.

The evaluation procedure hierarchy is currently classified under *Procedure by method*, with many immediate children as follows:

- *Procedure by method (procedure)*
  - Evaluation procedure (procedure); some children include:
    - Imaging (procedure)
    - Measurement procedure (procedure)
    - Physical examination assessment (procedure)
Evaluation procedures can be defined by a Method (attribute) of Evaluation - action (qualifier value).

Subtypes of Evaluation-action (qualifier value) include:

- Examination - action (qualifier value)
- Imaging - action (qualifier value)
- Measurement - action (qualifier value)
- Monitoring - action (qualifier value)
- Spectroscopy - action (qualifier value)

Reason for procedure

In general, the reason for ordering a procedure should not be precoordinated within the procedure concept, i.e. it should not constrain the reporting of results. The reason that a procedure is ordered may influence the interpretation of the results but usually does not influence the way the procedure is performed.

Unacceptable example,

- Computed tomography angiography of chest with contrast for evaluation of pulmonary embolus (procedure)

Acceptable example including reason for procedure,

- 66596009 | Diagnostic radiography for foreign body detection and localization (procedure)

Study

Procedures with the word study are unacceptable. They are ambiguous, as they imply context beyond the execution of the procedure.
Primary vs secondary procedures

The meaning of primary and secondary, when describing a procedure, is open to interpretation. Consequently, the concepts will be inactivated.

The interpretation of primary may be:

- Not ever done before at this site
- The first of multiple procedures, with two sub-meanings:
  - The first of planned multiple procedures, whether the plan is carried out or not
  - The first of multiple procedures that were not planned or foreseen, i.e. it is only the first of multiple procedures in retrospect

Examples of unacceptable descriptions,

- Primary anterior decompression of cervical spinal cord (procedure)
- Primary anterior excision of cervical intervertebral disc (procedure)
- Primary arthrodesis of interphalangeal joint of toe (procedure)
- Primary anterolateral excision of thoracic intervertebral disc (procedure)

Centesis

Centesis may be defined as the act of puncturing a body cavity or space with a hollow needle and drawing out fluid. Each centesis procedure involves both a puncture action and a needle aspiration action. It is correct to have two relationship groups for centesis procedures.

One group has a Method (attribute) of Puncture - action (qualifier value) and a Procedure site - Direct (attribute) of the structure being punctured.

For example,

- 91602002 Thoracentesis (procedure) has Procedure site - Direct (attribute) of Pleural membrane structure (body structure)

The second group has a Method (attribute) of Aspiration - action (qualifier value) and a Procedure site - Indirect (attribute) of the space being aspirated.

For example,

- 91602002 Thoracentesis (procedure) has Procedure site - Indirect (attribute) of Pleural cavity structure (body structure)

Division, lysis, transection, bisection

Division and lysis

Division - action (qualifier value) is a subtype of Incision - action (qualifier value). This does not mean that all procedures, that include the word division, should necessarily be modeled with Method (attribute) of Division - action (qualifier value), like those where the division is accomplished using blunt dissection, not incision.

For example,
Division of adhesion concepts, like 173269002 Division of adhesions of lip (procedure), should be modeled the same as lysis of adhesion concepts, like 45602008 Lysis of adhesions of peritoneum (procedure). Both use Dissection - action (qualifier value). Adhesions are broken down by blunt dissection, often without incising them. This does not exclude procedures that may also involve division by incision.

The preferred name of division of adhesions concepts can be changed to lysis of adhesions for consistency. The use of lysis of adhesions also helps with correct modeling and avoidance of interpreting divisions as necessarily being kinds of incision.

Transection and bisection
Transection is defined as a division across the longitudinal axis of a structure by cutting. Bisection is defined as division into two parts by cutting. Transection - action (qualifier value) is a subtype of Bisection - action (qualifier value), which is a subtype of Division - action (qualifier value) and Incision - action (qualifier value).

For example,

- 53176004 Transection of muscle of eye (procedure)
- 60158005 Bilateral bisection of ovary (procedure)

Encounter
An encounter is defined as an in-person meeting between a patient and a healthcare provider for the purpose of the provision of healthcare services to the patient. 308335008 Patient encounter procedure (procedure) is a subtype of Procedure (procedure).

For example,

- 185349003 Encounter for check up (procedure)

An indirect encounter occurs without a face-to-face meeting. 185316007 Indirect encounter (procedure) is a subtype of 308335008 Patient encounter procedure (procedure).

For example,

- 386473003 Telephone follow-up (procedure)

Endoscopy vs. endoscopic procedure
Endoscopic procedures are distinguished from endoscopy procedures. The distinction depends on the Action (qualifier value) of the Method (attribute).

In an endoscopy, the Method is Inspection - action (qualifier value). For these procedures, Endoscope, device (physical object) is the value for Using device (attribute).

For example,

- 427595003 Capsule endoscopy (procedure) has the Relationship group
  - Using device, Endoscope, device (physical object)
  - Procedure site, Direct, Gastrointestinal tract structure (body structure)
  - Method, Inspection - action (qualifier value)

In an endoscopic procedure, the Method (attribute) has some other action. It is accomplished by gaining access to the procedure site via an endoscope. For these procedures Endoscope device (physical object) is the value for Using Access Device (attribute). This specifies that the endoscope is used to access the site.

For example,

- 53767003 Endoscopic biopsy (procedure) has the Relationship group
  - Using access device, Endoscope, device (physical object)
  - Method, Biopsy, action (qualifier value)
Excision, incision, biopsy

Excision, incision, and biopsy may be difficult to interpret. They are organized according to the following general structure.

Excision

Organ excision. Any excisional act involving the organ; usually (organ)-ectomy, or similar, is a synonym. Organ excision, itself, does not specify whether it is complete or partial, nor does it specify what is excised.

For example,
- 23968004 | Excision of colon (procedure) | or one of the synonyms, Colectomy

Complete or total excision

Concepts may include complete or total to indicate complete removal or excision of the organ.

For example,
- 63016009 | Total resection of urinary bladder (procedure) | with the synonyms Complete cystectomy, Total excision of bladder, and etc

Partial excision

Concepts may include partial to indicate removal or excision of part of the organ. Specifying partial excision does not differentiate between a partial excision of or from the organ.

For example,
- 708929007 | Laparoscopic partial excision of kidney using robotic assistance (procedure) | or one of the synonyms, Partial nephrectomy, laparoscopic with robot assistance

Lesion or tissue

Concepts may indicate removal of a lesion or tissue; excision of a lesion or tissue from an organ may be complete or partial.

For example,
- 72106008 | Excision of lesion of liver (procedure) |
- 69031006 | Excision of breast tissue (procedure) |

Lesion modeling

The word lesion can be used to refer to both structural and functional abnormalities. If a procedure (or disorder) refers to a lesion in a way that makes it clear that it is a generic term for a structural abnormality, then the correct modeling approach is to use Procedure morphology (attribute) for procedures or [Associated morphology (attribute), Morphologically abnormal structure (morphologic abnormality) for disorders].

Excision(al) biopsy

Excisional biopsy of organ generally means that tissue or a lesion or suspected lesion is necessarily entirely excised, not the entire organ. It is a partial excision of (from) the organ. This is true even when small polyps are removed.

For example,
- 116237003 | Excisional biopsy of lesion of rectum by transanal approach (procedure) |

Incision

An organ incision is any incisional act involving the organ; usually (organ)-otomy, or similar, is a synonym

For example,
- 45558009 | Incision of lung (procedure) | or the synonym, pneumonotomy

Any incision procedure that does not necessarily involve division (as opposed to ordinarily does not involve division) remains primitive without an available negation operator.
Incisional biopsy
Incisional biopsy of organ; incisional biopsy of lesion of organ; usually with open approach. Incisional biopsy of [organ] necessarily implies incision and removal of a lesion, and is by definition a *partial excision*, since the site is the organ, and an excision is done, but the entire lesion is not necessarily removed.

For example,

- 237378001 [Incisional biopsy of breast (procedure)]

### Modeling biopsies

Biopsies, like other removal procedures, may have two direct objects, the **morphology** and the **site**. It is permissible to use Procedure site - Direct (attribute) for biopsies, even if subtypes might have a direct object that is a morphology.

### Grafting

Although the use of terminology may vary across specialties, in general, grafting is where tissue is completely separated from its source of origin or donor, without its own blood supply, then affixed to a recipient site. The recipient site provides the vascularity.

**Graft of skin is**

- A section of skin with variable size, thickness, and origin
- Completely detached from its original site and moved to cover the area to be repaired without the benefit of any blood supply

For grafting, the recipient site is represented with the Procedure site - Direct (attribute), and the graft is represented with the Direct substance (attribute).

Fixation or attachment of tissue involves skin, bone, cartilage, or fat, rather than whole organs. The term can also be used for fixation or attachment of synthetic materials (e.g., a bioengineered skin graft is a manufactured skin graft grown in the laboratory from the patient’s own cells, other allogeneic or xenogeneic sources, and/or synthetic materials; for example, silicone graft).

For example,

- 783285007 [Full thickness graft of skin to skin of neck (procedure)]
  - Proximal primitive Is a (attribute) value is 71388002 Procedure (procedure)
  - A single relationship group consists of:
    - 260686004 [Method (attribute)]= 129407005 [Grafting - action (qualifier value)]
    - 405813000 [Procedure site - Direct (attribute)]= 43081002 [Skin structure of neck (body structure)]
    - 311501000 [Direct substance (attribute)]= 782792007 [Full thickness graft of skin (substance)]
      - Values for direct substance should be from the 420934007 [Graft of skin (substance)] hierarchy that include the origin of the material in the description.

⚠️ *Skin flaps* are under review and not included here
Imaging guidance

Imaging guidance can be modeled using the Has intent (attribute). The concept 429892002 |Guidance intent (qualifier value)|, a child of 363675004 |Intents (nature of procedure values) (qualifier value)|, is the value for the Has intent (attribute) for imaging-guided procedures.

For example,

- 432666003 |Biopsy of brain using computed tomography guidance (procedure)| has two relationship groups:
  - Method (attribute) of Biopsy - action (qualifier value)
  - Procedure Site - Direct (attribute) of Brain structure (body structure)
  - Procedure Site - Direct (attribute) of Brain structure (body structure)
  - Method (attribute) of Computed tomography imaging - action (qualifier value)
  - Has intent (attribute) of Guidance intent (qualifier value)

**Biopsy of brain using CT guidance**

432666003 |Biopsy of brain using computed tomography guidance (procedure)| is subsumed by 702707005 |Biopsy of head (procedure)| and by 34227000 |Computerized axial tomography of brain (procedure)|.

Fluoroscopic guidance

*X using fluoroscopic guidance (procedure)* is a subtype of Fluoroscopy (procedure).

For example,

- 710291004 |Endoscopy using fluoroscopic guidance (procedure)| has the following relationship groups:
  - Using device (attribute) of Endoscopic device (physical object)
  - Method (attribute) of Inspection - action (qualifier value)
  - Method (attribute) of Fluoroscopic imaging - action (qualifier value)
  - Has intent (attribute) of Guidance intent (qualifier value)

**See also Clinical imaging procedure naming conventions section**

Immunization and vaccination

Immunization may be active (introduction of a vaccine) or passive (introduction of immunoglobulin/antibodies). A vaccine is a substance that can induce active immunity. Vaccination, by definition, is the introduction of a vaccine, and is, therefore, synonymous with active immunization. Some descriptions include the word vaccination, where it is clear that vaccination is intended. Other descriptions have preferred terms with the word vaccination, and synonyms with the word immunization, to include both active and passive immunization.

For example,

- 38598009 |Measles-mumps-rubella vaccination (procedure)| has vaccination in all descriptions
- 86198006 |Influenza vaccination (procedure)| has the synonym, influenza immunization

Regimes and therapies

A regime/therapy is a set, sequence, or group of procedures. 243120004 |Regimes and therapies (regime/therapy)| is a subtype of Procedure (procedure). As a subtype of procedure, Regimes and therapies have the same attributes and use the same model as general procedures.
Regime and therapies are either repeated multiple times, over an extended period of time.

For example,

- 716872004 | Antineoplastic chemotherapy regimen (regime/therapy), This regime/therapy might include individual instances of administration of chemotherapy agents; the instances are at separate times, over a predetermined or planned period of time.
- 229586001 | Rest, ice, compression and elevation treatment program (regime/therapy), This regime/therapy refers to repeated rest, ice, compression and elevation (RICE) for an indefinite period of time.

Regime and therapies are focused on a single purpose but do not have any single sub-procedure as a necessary part.

For example,

- 385695003 | Cast care (regime/therapy), The sub-procedures are all done for the purpose of properly monitoring and maintaining an orthopedic cast, but the sub-procedures may vary from one cast, patient, or healthcare setting to the next. Sub-procedures may include inspecting the cast, checking the skin, reinforcing padding, etc. There is not a single sub-procedure as a necessary part, although the purpose of the sub-procedures is to take care of a cast.
- It is possible to have a regime/therapy as an instance of care. An instance of cast care could be the specific care for Mr. Smith’s cast on the morning of April 23rd, consisting of the set of procedures: examining the cast; examining his arm; asking about his symptoms; and cleaning the skin.

**Has focus**

*Regime/therapy* may be the value for the *Has focus (attribute)*.

For example,

- 385978009 | Cardiac rehabilitation assessment (procedure) has a Has focus (attribute) of Cardiac rehabilitation (regime/therapy)

**Skeletal system**

Since the skeletal system includes bones and cartilage, it is possible to have a procedure on the skeletal system, i.e. on cartilage, that is not a procedure on bone.

For example,

- 77825002 | Division of cartilage of wrist (procedure) is a procedure on the skeletal system (procedure)

**Skeletal system subdivision**

SNOMED CT considers the *skeletal system subdivision* part of the entire bone (system). This may change if there are procedures on cartilaginous skeleton that involve skeletal system subdivisions.

**Osteotomy**

*Osteotomy* is defined as *cutting into or through a bone*; there are 3 meanings in SNOMED CT:

1. Cutting into a bone, regardless of whether the bone is divided (incision, general meaning). Model using Method, Incision - action (qualifier value), and Procedure site - Direct (attribute), bone structure (or subtypes).
   
   **For example,**

   - 118483001 | Incision of rib (procedure)

2. Cutting through a bone and dividing it (division by cutting). Model using Method, Division - action (qualifier value), and Procedure site - Direct (attribute), bone structure (or subtypes).

   **For example,**
3. Cutting into a bone without cutting through it and therefore without dividing it (incision without division).

This is unnecessary; procedures that do not explicitly involve division are modeled simply as *Incision*.

Reduction and fixation of fractures

*Reduction and fixation* has two actions by two different means; open reduction of a fracture and insertion of an orthopedic fixation device. This provides an opportunity for general concept inclusion axioms (GCIs) in order to fully represent the meanings without heavy postcoordination modeling. *Open reduction of a fracture* necessarily involves open manipulation of the fracture and *internal fixation of a fracture* necessarily involves the insertion of an orthopedic internal fixation device.

For example,

- 74011006 | Open reduction of fracture of tibia and fibula with internal fixation (procedure)

### Surgical procedures

A *surgical procedure* is defined as a procedure that involves intentional non-transient alteration of structures of the body, and/or a procedure that necessarily involves cutting into the body. This definition includes all procedures defined by *Method* (attribute) with Surgical action (qualifier value).

SNOMED CT classifies concepts as surgical procedures if their methods are *surgical actions* based on the action hierarchy. The surgical action hierarchy distinguishes surgical from non-surgical actions based on the definition above. Note the *Or* in the sentence; actions that do not involve cutting or incision, but do involve the intentional non-transient alteration of anatomy, are still surgical.

---

**Operation**

In SNOMED CT, *operation* is synonymous with surgical procedure.

Surgical procedures are not defined simply as procedures done by a surgeon (despite some dictionary definitions). Surgeons can perform many non-surgical actions and surgical procedures need not necessarily be performed by a surgeon, i.e. if a non-surgeon performs a surgical procedure, it is still a surgical procedure.

---

**Medical procedure**

The use of the term *medical procedure* is deprecated, i.e. not recommended, because it lacks reproducible meaning. It might be defined as *a procedure done by a physician*, but even that is deprecated, because it is provider-specific.

---

**Revision**

A *revision procedure* is not a subtype of the original procedure. Revision procedure concepts should be in the 118635009 | Revision (procedure) | sub-hierarchy.

For example,

- 171839006 | Re-release of carpal tunnel (procedure) | is modeled as follows:
Surgical repair

The definition of *surgical repair* is restoring, to the extent possible, an anatomical structure, using a surgical action. *Repair* is an objective or intended accomplishment, not a means (e.g. suturing, transplanting, etc.) nor a need (e.g. normal functioning, cosmetic appearance, pain relief, etc.).

Surgery that restores structure is usually intended to restore function and appearance. Restoring function, however, is not necessary for a procedure to be considered a repair. It is also possible for surgery to restore function without restoring structure (e.g. surgery to attach a prosthetic limb after amputation). This type of surgery would not be strictly categorized as a repair.

The *Method* (attribute) is used to model both the objective of a procedure and the means used to accomplish it. If a procedure requires both a repair action and another type of action, then two relationship groups should be used.

**Fistula**

*Closure* action is a kind of *repair* action. All fistula closures use the *closure* action and are classified as kinds of repair procedures.

For example,
• 79433000 | Closure of colon fistula (procedure) | has Method (attribute), Closure - action (qualifier value) with a parent, Repair of colon (procedure)

Plastic repair
Surgery that accomplishes a repair (a structural restoration) often uses the suffix -plasty. The term plastic repair is also used. In order to avoid redundancy, the following terms are used:

• Prosthetic repair, using external (non-body) materials
• Plastic repair, reshaping the body

- ► Plasty
The suffix -plasty is widely used in concepts that apply to prosthetic repairs (e.g. total hip arthroplasty). So -plasty may refer to any general repair (prosthetic, plastic, or other) and not just plastic repairs.

Surgical vs. non-surgical
As mentioned in the initial Surgical procedures page, the definition of surgical procedure includes intentional non-transient alteration of structures of the body and/or necessarily involves cutting into the body.

Non-surgical actions do not significantly or non-transiently alter anatomy and do not necessarily involve cutting or incision.

For example,

• Fine needle biopsy (procedure) or brush biopsy (procedure)
• Phlebotomy, a synonym for venipuncture for blood test (procedure)
• Aspiration (procedure)
• Closed reduction of dislocation (procedure)

Closed procedure naming

• The general pattern <open, closed> <procedure> is accepted
• When a procedure is specified as closed, the closed procedure should be fully described, e.g. fine needle biopsy, endoscopic, etc.

Under revision

48635004 | Fine needle biopsy (procedure) | could be viewed as a kind of centesis, but the former is non-surgical and the latter is surgical. Sampling - action (qualifier value), in general, is not necessarily a surgical action. If sampling involves the surgical removal of part of something, then Surgical biopsy (procedure) should be the action.

Qualifier Value

<table>
<thead>
<tr>
<th>Definition</th>
<th>Examples</th>
</tr>
</thead>
</table>
| Qualifier values include a wide range of concepts that provide attribute values used in the definitions of other concepts. These values can also be used in expressions to refine the meaning of a concept, or in the appropriate fields of a health record to add additional information. | • Action, Numbers, Clinical specialty, Context values, Mode of transmission, Type of diagnosis, Scale type, Sport, Technique, Time frame, World languages
• There are also many Qualifier value subtypes relating to the pharmaceutical realm: Additional dosage instructions, Basic dose form, Route of administration value, etc. |

The 362981000 | Qualifier value (qualifier value) | hierarchy contains concepts used as the target value of an attribute in a defining relationship.

For example,

• 18639004 | Left kidney structure (body structure) | has a Laterality (attribute) of Left (qualifier value)
The use of qualifiers varies greatly according to the domain to which they are applied. Thus, guidance in their use is often found within the guidance of the respective domain. So for the above example, information regarding the laterality/left attribute-value pair, see the Laterality guidance found within the Body structure domain. The range of values for a particular attribute is provided in the specific concept model of the domain. For further information on the range of values for a specific domain, see the different content types and rules in the MRCM maintenance tool at https://browser.ihtsdotools.org/mrcm.

Specific information on a few select subhierarchies can be found below:

### Disposition

The 726711005 | Disposition (disposition) | hierarchy is required to support the remodeling of the 105590001 | Substance (substance) | hierarchy. These concepts are used as the attribute values for the 726542003 | Has disposition (attribute) |. The (disposition) semantic tag is used to differentiate concepts in this hierarchy from similar concepts in other hierarchies.

<table>
<thead>
<tr>
<th>FSN</th>
<th>X (disposition)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>For example,</td>
</tr>
<tr>
<td></td>
<td>• Coagulation factor inhibitor (disposition)</td>
</tr>
<tr>
<td></td>
<td>• Acute phase reactant (disposition)</td>
</tr>
<tr>
<td></td>
<td>• Human immunodeficiency virus fusion inhibitor (disposition)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>PT</th>
<th>X</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>For example,</td>
</tr>
<tr>
<td></td>
<td>• Coagulation factor inhibitor</td>
</tr>
<tr>
<td></td>
<td>• Acute phase reactant</td>
</tr>
<tr>
<td></td>
<td>• HIV fusion inhibitor</td>
</tr>
</tbody>
</table>

**Modeling**

*Techniques*, as qualifier values, should include the word *technique* in their FSNs.

For example,

- 702658000 | Microbial culture technique (qualifier value) |

### International System of Units - derived unit of volume

The 282115005 | International System of Units-derived unit of volume (qualifier value) | hierarchy contains concepts representing metric units of volume.

<table>
<thead>
<tr>
<th>FSN</th>
<th>X metric unit of volume</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Lower case with case sensitivity, ci</td>
</tr>
</tbody>
</table>
### International System of Units - unit of mass

The [258681007 |International System of Units unit of mass (qualifier value)] hierarchy contains concepts representing metric units of mass.

<table>
<thead>
<tr>
<th>Concepts representing Unit of Mass</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>FSN</strong></td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td><strong>PT</strong></td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td><strong>SYN</strong></td>
</tr>
<tr>
<td></td>
</tr>
</tbody>
</table>

### Number

Following the deployment of the concrete domain functionality in SNOMED CT, concepts in the [260299005 |Number (qualifier value)] hierarchy are no longer necessary, and plans have been made for their inactivation in January 2023.

Concepts that describe numeric values have been added as descendants of [260299005 |Number (qualifier value)]. While the FSN and PT for these concepts use numeric characters, the concepts represent descriptions of a number and are not actually numeric values. To reinforce this, a synonym with the textual description of the number is created for each concept.

For example,

- |Zero point two| is the synonym for [732349004 |0.2 (qualifier value)]
Concepts representing numbers

<table>
<thead>
<tr>
<th>FSN</th>
<th>X (qualifier value)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Trailing zeros are not allowed (e.g. 10, not 10.0)</td>
</tr>
<tr>
<td></td>
<td>Preceding zeros are required (e.g. 0.5, not .5)</td>
</tr>
<tr>
<td></td>
<td>For example,</td>
</tr>
<tr>
<td></td>
<td>• 25 (qualifier value)</td>
</tr>
<tr>
<td></td>
<td>• 37.5 (qualifier value)</td>
</tr>
<tr>
<td></td>
<td>• 125 (qualifier value)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>PT</th>
<th>X</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Trailing zeros are not allowed (e.g. 10, not 10.0)</td>
</tr>
<tr>
<td></td>
<td>Preceding zeros are required (e.g. 0.5, not .5)</td>
</tr>
<tr>
<td></td>
<td>For example,</td>
</tr>
<tr>
<td></td>
<td>• 25</td>
</tr>
<tr>
<td></td>
<td>• 37.5</td>
</tr>
<tr>
<td></td>
<td>• 125</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>SYN</th>
<th>A synonym representing the concept as a textual description is required</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Synonyms are not case sensitive. No commas or other punctuation is allowed. An exception is the hyphen</td>
</tr>
<tr>
<td></td>
<td>For example,</td>
</tr>
<tr>
<td></td>
<td>• Twenty-five</td>
</tr>
<tr>
<td></td>
<td>• Thirty-seven point five</td>
</tr>
<tr>
<td></td>
<td>• One hundred and twenty-five</td>
</tr>
</tbody>
</table>

Pharmaceutical dose form subhierarchy

The 736542009 |Pharmaceutical dose form (dose form)| subhierarchy of 362981000 |Qualifier value (qualifier value)| contains concepts which support the Medicinal product model.

For these concept model domains that support the drug model, see the attribute and range tables at the Pharmaceutical and Biologic Product Attributes Summary page.

Please see the SNOMED CT Pharmaceutical Dose Form Editorial Guide for information on the following qualifier value subhierarchies that support the drug model.

- 736478001 |Basic dose form (basic dose form)|
- 736665006 |Dose form administration method (administration method)|
- 736479009 |Dose form intended site (intended site)|
- 736480007 |Dose form release characteristic (release characteristic)|
- 736477006 |Dose form transformation (transformation)|
- 736471007 |State of matter (state of matter)|

Type of drug preparation

Descendants of 105904009 |Type of drug preparation (qualifier value)| do not meet the criteria to be considered Pharmaceutical dose forms. This subhierarchy will be retained as a primitive subhierarchy until such time that use cases and/or detailed requirements are known. Requests for addition of new concepts or modification of existing concepts will be evaluated on a case-by-case basis.
Unit of presentation

A unit of presentation represents a qualitative concept that describes a countable entity in which the clinical drug is presented (e.g. tablet, capsule) or in which it is bounded (vial, ampule). The 732935002 | Unit of presentation (unit of presentation) hierarchy supports harmonization between SNOMED CT’s Drug Concept Model and the International Organization for Standardization’s Identification of Medicinal Products (IDMP) standards for product strength.

Out of Scope

- Concepts representing proprietary dose forms
- Concepts that contain modifiers, e.g. hard capsule, capsule for inhalation

⚠️ Unit dose

The Unit dose (qualifier value) is unacceptable for representing unit of presentation.

<table>
<thead>
<tr>
<th>Concept descriptions representing Unit of Presentation</th>
</tr>
</thead>
<tbody>
<tr>
<td>FSN</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td></td>
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<td></td>
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<tr>
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</tbody>
</table>

<table>
<thead>
<tr>
<th>PT</th>
<th>X</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>For example,</td>
</tr>
<tr>
<td></td>
<td>• Actuation</td>
</tr>
<tr>
<td></td>
<td>• Capsule</td>
</tr>
<tr>
<td></td>
<td>• Suppository</td>
</tr>
<tr>
<td></td>
<td>• Tablet</td>
</tr>
</tbody>
</table>

| SYN  | Synonyms are not allowed |

Record Artifact

<table>
<thead>
<tr>
<th>Definition</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Clinical documents, or parts thereof</td>
<td>• 422813005</td>
</tr>
<tr>
<td></td>
<td>• 416575001</td>
</tr>
</tbody>
</table>

A record artifact is an entity that is created by a person or persons for the purpose of providing other people with information about events or states of affairs.

In general, a record is virtual, that is, it is independent of its particular physical instantiation/s. It consists of information elements (usually words, phrases and sentences, but also numbers, graphs, and other information elements).

Record artifacts need not be complete reports or records. They can be parts of a larger Record artifact.

For example,
• **A184225006 Computer record of patient (record artifact)** is a Record artifact that also may contain other Record artifacts in the form of individual documents or reports, e.g. **726738003 Cytology report (record artifact)**. These may, in turn, contain more finely granular Record artifacts, such as sections, and even section headers e.g. **422813005 Document section (record artifact)**.

### Situation with Explicit Context

<table>
<thead>
<tr>
<th>Definition</th>
<th>Examples</th>
</tr>
</thead>
</table>
| Concepts that include context information; a subtype of the situation to which it applies, with an attribute associating it with the relevant clinical finding or procedure | **407565004 Angiotensin II receptor antagonist not tolerated (situation)**  
**417886001 Treatment adjusted per protocol (situation)** |

### Variable meanings according to context

Depending on context, concepts can be used in many different ways with various meanings.

**A disorder concept** can represent:

- Possible diagnosis or part of a differential diagnosis
- Diagnosis applied to a family member or some other contact person
- Diagnosis explicitly excluded
- Diagnosis, now known to be incorrect, but which was the basis for a particular course of treatment
- Absent feature of a related disorder
- Diagnosis that the patient believes or fears they have

**A procedure concept** can represent:

- Requested, recommended or planned procedure
- Procedure for which consent has been given or withheld
- Procedure that is contraindicated
- Procedure that has been canceled or postponed
- Procedure for which follow up is now being arranged
- Procedure which caused a complication

**A symptom concept** can represent:

- Confirmed absence of a symptom
- Symptom deduced and reported by a third party as a witness of a clinical event
- Inability or failure to obtain information about a symptom
- Symptom which the patient is advised to respond to in a particular manner

**A finding concept** can represent:

- Absence of a finding
- Inability or failure to check for a finding
- Finding which, if present, is to trigger a particular change in clinical management
- Finding which is the goal or target of a treatment

**A product concept** can represent:

- Allergy or other contraindication to a product
- Assertion that a product caused a particular side effect
- Various therapeutic activities of a product
- Instructions given to a patient for use of a non-prescription medication
- Clinical authorization of a prescription
- Issuing of a prescription for a course of treatment
- Supply (dispensing) of a specified quantity of a product
- Administration of a single dose of a product
- Change of a product dosage
• Discontinuation of a product
• Specialist’s recommendation to use a particular product, if certain circumstances apply

Situation with Explicit Context Attributes Summary

When authoring in this domain, these are the approved attributes and allowable ranges. They are from the Human Readable Concept Model (HRCM).

<table>
<thead>
<tr>
<th>Domain Information for</th>
<th>Situation with explicit context (situation)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Domain Constraint</td>
<td>&lt;&lt; 243796009</td>
</tr>
<tr>
<td>Parent Domain</td>
<td>-</td>
</tr>
<tr>
<td>Proximal Primitive Constraint</td>
<td>&lt;&lt; 243796009</td>
</tr>
<tr>
<td>Proximal Primitive Refinement</td>
<td>-</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Author View of Attributes and Ranges for</th>
<th>Situation with explicit context (situation)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Attribute</td>
<td>Grouped</td>
</tr>
<tr>
<td>408732007 [Subject relationship context (attribute)]</td>
<td>1</td>
</tr>
<tr>
<td>408731000 [Temporal context (attribute)]</td>
<td>1</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Author View of Attributes and Ranges for</th>
<th>Finding with explicit context (situation)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Attribute</td>
<td>Grouped</td>
</tr>
<tr>
<td>246090004 [Associated finding (attribute)]</td>
<td>1</td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td>408729009 [Finding context (attribute)]</td>
<td>1</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Author View of Attributes and Ranges for</th>
<th>Procedure with explicit context (situation)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Attribute</td>
<td>Grouped</td>
</tr>
<tr>
<td>363589002 [Associated procedure (attribute)]</td>
<td>1</td>
</tr>
<tr>
<td>408730004 [Procedure context (attribute)]</td>
<td>1</td>
</tr>
</tbody>
</table>
Situation with Explicit Context Defining Attributes

The following defining attributes correspond to the Situation with Explicit Context Attributes Summary table. Associated finding and Finding context are used with Findings with Explicit Context.

Associated finding

This attribute links concepts in the Situation with explicit context hierarchy to their related Clinical finding or Event. It specifies the Clinical finding or Event concept whose context is being modified.

When Associated finding is used in post-coordinated expressions, its range is broader than when used in pre-coordinated content. Associated finding should not reference concepts that already have pre-coordinated context.

For example,

- 443999008 | Risk of exposure to communicable disease (situation) | with Associated finding, Exposure to communicable disease (event)

For example, to create the concept, History of thyroid disease in father,

- Subject relationship context (attribute) with the value, father (person)
- Associated finding (attribute), with the value, Disorder of thyroid gland (disorder)

Incorrect example,

- Using Family history with explicit context (situation),
  - Subject relationship context (attribute) with the value, father (person)
  - Associated finding with value, Family history: Thyroid disorder (situation)

Finding context

This attribute represents a situation in which a Clinical finding or Event is known or unknown. If known, whether it is present, absent, or uncertain (possible). It also represents that the finding is not actual, but anticipated or possible in the future.

For example,

- 161922009 | No cough (situation) | with Associated finding, Cough (finding) and Finding context, Known absent (qualifier value)

Subject relationship context and Temporal context are used with Situations, Findings, and Procedures with Explicit Context.

Subject relationship context

This attribute is used to specify the subject of the Clinical finding or Procedure being recorded, in relation to the subject of the record.

For example,

- 161077003 | Father smokes (situation) | with Associated finding, Smoker (finding) and Subject relationship context, Father of subject (person)

Temporal context

This attribute indicates the time of the procedure or finding. It may be actual, i.e. occurred in the present, in the past, at a specified time; or in the future, i.e. it is planned or expected. The most general value is simply Current or past (actual), meaning that the concept was actual (not planned or expected), but not specifying anything further about the time. The word specified in the Temporal context means that there is a date or time stamp associated with the concept in the record. The date and/or time is a point and/or interval, that applies to the concept.

For example,
• 161550001 | History of hematuria (situation) | with Associated finding, Blood in urine (finding) and Temporal context, In the past (qualifier value)

Associated procedure and Procedure context are used with Procedures with Explicit Context.

Associated procedure
This attribute links concepts in the Situation with explicit context hierarchy to concepts in the Procedure hierarchy for which there is additional context.

For example,
• 183976008 | Operative procedure planned (situation) | with Associated procedure, Surgical procedure (procedure)

Procedure context
This attribute indicates the degree of completion, or status, of a Procedure, as well as its possible future states, prior to it being initiated or completed.

For example,
• 183976008 | Operative procedure planned (situation) | with Procedure context, Planned (qualifier value)

Situation with Explicit Context Naming Conventions
For information on precoordinated naming patterns that have been reviewed or are currently in review, see Pre-coordination Naming Patterns project. Unreviewed patterns for the Situation with explicit context hierarchy can be found here. New content should conform with the naming patterns; however, legacy content may not.

For example,
Acceptable naming pattern
FSN: <procedure> declined (situation)
PT: <procedure> declined
SYN: <procedure> refused (This is optional.)
• 736013005 | Body weight measurement declined (situation)|

Unacceptable naming patterns
The following naming patterns are no longer accepted for addition to the International Edition:
• Procedure offered
• Procedure not offered
• Procedure done
• Procedure not done

Note that the 385658003 | Done (qualifier value) | (a descendent of 410523001 | Post-starting action status (qualifier value)) remains in use as the target value of the 408730004 | Procedure context (attribute) | in History of <procedure> concepts.

Not every naming pattern is found in the Pre-coordination Naming Pattern project. Some naming patterns can be prescribed in templates. Others can come from trackers or fast track documents, such as the examples below.

No known allergy
The pattern is:
716186003 | No known allergy (situation)

FSN: No known allergy (situation)
PT: No known allergy
SYN: NKA - No known allergy

No known X allergy (situation)
For example, 428197003 | No known insect allergy (situation)
FSN: No known insect allergy (situation)
PT: No known insect allergy

Situation with Explicit Context Modeling
SNOMED CT contains concepts that include context information, and concepts that are regarded as context-free. A concept includes context information if the name of the concept explicitly represents information that might otherwise be represented by another less context-rich concept in a particular place within an electronic health record or EHR.

In SNOMED CT, context describes the effects of embedding a concept in a clinical situation, i.e. when it is used in an EHR.

For example,
- When the concept 22298006 | Myocardial infarction (disorder) is used in an EHR, it takes on a specific contextualized meaning. The meaning might be an assertion by the person entering the information, that on a given date, the patient was diagnosed with a myocardial infarction. Or, it may be used to document a complication of smoking, a protocol for chest pain, a medication contraindication, a part of a patient's medical history, a possible diagnosis justifying a diagnostic test, a diagnosis excluded by a diagnostic test, a patient's family history, etc.
- The concept for breast cancer, 254837009 | Malignant neoplasm of breast (disorder), might be used to indicate either a current diagnosis of breast cancer, a family history of breast cancer, or a past history of breast cancer. Each of these three meanings differs in regard to the context in which breast cancer is described.
  - Current diagnosis of breast cancer indicates that the breast cancer is present now, and in this patient.
  - Family history of breast cancer refers to breast cancer occurring in a family member of a patient.
  - Past history of breast cancer indicates that the breast cancer occurred in the patient, at some time in the past, and it is not necessarily present now.

Not only are the differences significant relative to a patient's health record, but they are also important to population-based data retrieval; e.g. it is incorrect to retrieve those who have a family history of breast cancer when searching for patients with a diagnosis of breast cancer.

Default context
When a SNOMED CT concept appears in an EHR without any explicitly stated context, that concept is considered to have a default context. However, the information in the health record structure or information model, can override the default context.

Default context for a Clinical finding concept implies that the finding is present (vs. being absent), that it applies to the subject of the record (the patient), and that it is current (or at a specified time in the past, linked to the concept).
Default context for a Procedure concept implies that the procedure was completed, that it was performed on the subject of the record (the patient), and that it was done at the present time (or at a specified time in the past, linked to the concept).

Explicit context
Concepts in the Situation hierarchy (given the appropriate record structure) have explicit context and can represent Clinical findings and Procedures that:

**Have not yet occurred**
For example,

- 165137000 | Endoscopy arranged (situation) |

**Refer to someone other than the patient**
For example,

- 160303001 | Family history: Diabetes mellitus (situation) |
- 395083002 | Discussed with next of kin (situation) |

**Have occurred at some time prior to the time of the current entry in the record**
For example,

- 161514008 | History of aortic aneurysm (situation) |

Attributes
These attributes are used to represent Clinical finding and Procedure concepts within the Situation hierarchy:

<table>
<thead>
<tr>
<th>Attributes</th>
<th>Clinical Finding</th>
<th>Procedure</th>
</tr>
</thead>
<tbody>
<tr>
<td>Associated finding</td>
<td>Associated finding</td>
<td>Associated procedure</td>
</tr>
<tr>
<td>Finding context</td>
<td>Finding context</td>
<td>Procedure context</td>
</tr>
<tr>
<td>Subject relationship context</td>
<td>Subject relationship context</td>
<td>Subject relationship context</td>
</tr>
<tr>
<td>Temporal context</td>
<td>Temporal context</td>
<td>Temporal context</td>
</tr>
</tbody>
</table>

Expressing context
Context typically alters the meaning of a concept, i.e. the resulting concept is no longer a subtype of the original concept.

**Precoordinated expression.** Clinical context is specified in the description and entered into a field in a patient's EHR.

For example,

- The precoordinated expression 266897007 | Family history: Myocardial infarction (situation) | might be put directly in a blank field in a patient's EHR. A family history of myocardial infarction is not a subtype of myocardial infarction, so family history modifies the context.
- The precoordinated expression 54355006 | Intracranial injury, without skull fracture (disorder) | might be put directly in a blank field in a patient's EHR. The disorder Intracranial injury, without skull fracture is not a subtype of skull fracture, so without modifies the context.

**Postcoordinated expression.** Clinical context is specified by combining concepts.

For example,

- 281666001 | Family history of disorder (situation) |, combined with 246090004 | Associated finding (attribute) | = 22298006 | Myocardial infarction (disorder) |. These two concepts indicate a family history of myocardial infarction.
Concept or expression in an EHR field. A concept is placed in a field with a predefined meaning in an electronic health record. The meaning is conveyed by the context in which it is recorded.

For example,

- **Hip replacement planned** might be represented as 397956004 | Prosthetic arthroplasty of the hip (procedure) within a section of a patient’s health record called Planned actions. A planned hip replacement is not a kind of hip replacement, so the Planned actions record section modifies the context.

- **2004005 Normal blood pressure (finding)** might be placed in a field labeled as Goal in a patient’s EHR. A goal of normal blood pressure is not a kind of Normal blood pressure (finding), so the Goal field in the EHR modifies context.

Elaboration: changing concept meaning

**Elaboration** in SNOMED CT refers to any addition to or change of the meaning of a concept that may be brought about when it is embedded in a clinical situation. Embedding a concept in a clinical situation may **elaborate** the semantic interpretation of a concept in one of the following ways:

1. **Subtype qualification**
2. **Axis modification**
3. **Affirmation or Negation**
4. **Combination**

**Subtype qualification**

A **subtype qualification** refines the meaning of a concept. **Subtype qualification** is elaboration that results in a concept that is a subtype of the original unelaborated focus concept. A focus concept is the part of a SNOMED CT expression that represents a clinical finding, observation, event, or procedure. It may be given context by a surrounding context wrapper and may be made more specific by a refinement.

For example,
A past history of replacement of the left hip may be represented by a SNOMED CT expression in which the focus concept, hip replacement, is refined by laterality, left and enclosed in a context wrapper representing past history.

Subtype qualification

Subtype qualification has also been called a qualifier (e.g. ENV136060, GEHR, CTV3) or a secondary status term (e.g. NHS Context of Care). In SNOMED CT, the term subtype expresses more clearly the distinctive property of a qualifier. This is helpful because the meaning of modify and qualify are synonymous in many dictionaries and by some International Organization of Standardization (ISO) authorities.

Axis modification

The attributes used to define situation concepts permit explicit (rather than default) representation of various contexts. These attributes can change the meaning of a Clinical finding or Procedure concept in a way that changes the hierarchy (or axis) of the concept from Clinical finding or Procedure to Situation with explicit context. The resulting modified meaning is not a subtype of the original meaning of the concept, and therefore the axis-modifying attributes are not used to qualify the concept, but instead are used to qualify a Situation concept.

For example,

- The concept 22298006 | Myocardial infarction (disorder) may be elaborated by including it in a clinical record specifying family history. A record of a family history of myocardial infarction does not imply that the patient has had any type of myocardial infarction. Therefore, family history changes the focus from the default context to a specified context.
- The concept 52734007 | Total replacement of hip (procedure) may be elaborated by stating that the procedure is planned for some future date. A record of planned total hip replacement does not imply that the patient has actually had a total hip replacement, i.e. it is not the default context for a procedure.
- The concept 167272007 | Urine protein test not done (situation) uses the context-modifying attribute Procedure context (attribute) and a value of Not done (qualifier value). This concept is not a subtype of 167271000 | Urine protein test (procedure), because its axis (hierarchy) is modified. Note that |<Procedure> not done| is no longer allowed. See the list disallowed naming patterns at Pre-coordination Naming Patterns Project.

Affirmation and Negation

Depending on perspective, affirmation and negation may simply be viewed as the inversion of meaning of an unelaborated concept that represents a Clinical finding. A concept may be stated in the negative in a clinical situation (e.g. meningism not present). This creates the potential for a concept to represent two meanings, one of which is the inverse of the other. However, the effects of negation on interpretation are far-reaching and distinct from other elaborations.

Negation, like axis modification, results in a concept that is not a subtype of the unelaborated concept. However, negation explicitly rules out the unelaborated concept.

For example,

- Family history of myocardial infarction does not imply that a patient had a myocardial infarction.
- No headache implies that patient has headache is untrue. A negative statement may expand further in the opposite direction of a positive statement. If headache is a subtype of pain then patient has headache implies patient has pain. However, patient has no headache does not imply patient has no pain. Conversely, patient has headache does not imply patient has occipital headache, but patient has no headache implies patient does not have occipital headache.
Negation

The representation of negation within SNOMED CT that arises from restrictions imposed by the existing description logics results in the hierarchy being inverted e.g., coronary heart disease not present is NOT properly a subtype of "Heart disease not present", which is clearly incorrect. An initial attempt was made to move negated content into the situation hierarchy so that the content remained available but SNOMED International recommends handling negation outside of SNOMED CT by the EHR vendor rather than try and represent it incorrectly within the terminology.

A concept may be stated to be possible in a clinical situation. Statements that explicitly indicate uncertainty can be considered in two possible ways:

- Somewhere between affirmation and negation
- As a type of elaboration

Combination

Two or more concepts may be embedded in a clinical situation in a way that links them together. Linkages may include:

- Simple combination of concepts
- Combination of a concept that is present and another that is absent

Context shift

Once a concept has context-shifted and become context-dependent, it should not be used in an expression that once again shifts context. In other words, when one context attribute is given an axis modifying value, the other context attributes are fixed.

For example,

- The model for 430679000 | Family history of diabetes mellitus type 2 (situation) | IS A Situation with explicit context (situation) with:
  - Subject relationship context of Person in family of subject (person)
  - Associated finding of Diabetes mellitus type 2 (disorder)
  - Finding context of Known present (qualifier value)
  - Temporal context of Current or past (actual) (qualifier value)

Even though the Family part of the concept results in an explicit axis shift of the Subject relationship context only, SNOMED CT requires default values for Finding context and Temporal context, rather than allowing them to be unspecified.

To negate a concept with Finding context (attribute) of Known present (qualifier value), the Finding context (attribute) should instead have a value of Known absent (qualifier value).

For example,

- The concept 160273004 | No family history: Hypertension (situation) | negates 160357008 | Family history: Hypertension (situation) by changing the value of Finding context (attribute) to Known absent (qualifier value) with Temporal Context (attribute) of All times past (qualifier value). The parent IS A Situation with explicit context (situation) with:
  - Temporal context of All times past (qualifier value)
  - Associated finding of Hypertensive disorder, systemic arterial (disorder)
  - Finding context of Known absent (qualifier value)
  - Subject Relationship Context of Person in family of subject (person)
SNOMED CT Model Component

<table>
<thead>
<tr>
<th>Definition</th>
<th>Types</th>
</tr>
</thead>
<tbody>
<tr>
<td>Concepts and attributes necessary to organize and structure SNOMED CT</td>
<td>900000000000442005</td>
</tr>
<tr>
<td>terminology and its derivatives</td>
<td>900000000000454005</td>
</tr>
<tr>
<td></td>
<td>106237007</td>
</tr>
<tr>
<td></td>
<td>370136006</td>
</tr>
</tbody>
</table>

SNOMED Model component module (metadata)

- SNOMED CT Model Component (metadata)
  - Core metadata concept (core metadata concept)
    - Case significance (core metadata concept)
    - Characteristic type (core metadata concept)
    - Definition status (core metadata concept)
    - Description type (core metadata concept)
    - Identifier scheme (core metadata concept)
    - Modifier (core metadata concept)
    - Module (core metadata concept)
  - Foundation metadata concept (foundation metadata concept)
    - Reference set (foundation metadata concept)
      - Reference set attribute (foundation metadata concept)
  - Linkage concept (linkage concept)
    - Attribute (attribute)
    - Link assertion (link assertion)
  - Namespace concept (namespace concept)
    - Core Namespace (namespace concept)
    - Extension Namespace (1000000) (namespace concept)
    - Extension Namespace (1000001) (namespace concept)
    - Extension Namespace (1000002) (namespace concept)
    - Extension Namespace (1000003) (namespace concept)
    - Extension Namespace (1000004) (namespace concept)

Core metadata concept

Subtypes of 900000000000442005 | Core metadata concept (core metadata concept) provide structural information required to support International Release data. This supporting information includes sets of enumerated values that apply to attributes of concepts, descriptions, and relationships.

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Foundation metadata concept

Subtypes of the \texttt{900000000000454005 \mid Foundation metadata concept (foundation metadata concept)} provide supporting metadata and structural information for derivative release structures including Reference Sets.

Linkage concept

A \texttt{106237007 \mid Linkage concept (linkage concept)} links two or more concepts to express compositional meanings. All concept codes that can be used as a \textit{Relationship Type} are included under Linkage concept (linkage concept). The Concept Model attributes are approved for use.

Linkage concept is a subtype of \texttt{900000000000441003 \mid SNOMED CT Model Component (metadata)}. The Linkage concept hierarchy has the subhierarchies:

- Attribute (attribute)
- Link assertion (link assertion)

Concepts in the Linkage concept subhierarchy are used to construct relationships between two SNOMED CT concepts; they indicate the relationship type between those concepts. Some attributes (relationship types) can be used to logically define a concept (defining attributes).

Unapproved attributes

\texttt{408739003 \mid Unapproved attribute (attribute)} is a subtype within this hierarchy with over a thousand descendants. Unapproved attributes in the SNOMED CT Concept model may be used to create post-coordinated expressions with caution. Use of unapproved attributes is neither supported by the MRCM nor recommended beyond intraorganizational use. Approved attributes are those that fit the MRCM for data sharing and interoperability between systems. Unapproved attributes are used for creating expressions within a single system for semantic matching of vendor terms (i.e. those that are used for mapping of interface terms, clinical decision support that use components within the expression, etc.).

Users should beware that unapproved attributes can potentially

- conflict with approved attributes if used without discretion
- change into an approved attribute if warranted by SNOMED International

Namespace concept

\texttt{370136006 \mid Namespace concept (namespace concept)} is a subtype of \texttt{900000000000441003 \mid SNOMED CT Model Component (metadata)}. Each of its subtypes has an integer term which is an assigned extension namespace identifier.
For more information

New namespace concepts are requested via email to info@snomed.org. A SNOMED International staff terminologist will add new extension namespace identifiers as requests are received. It is also necessary to change the Module ID per the following:

For further details, see Namespace identifiers on the SNOMED website at: https://www.snomed.org/snomed-ct/Use-SNOMED-CT.

<table>
<thead>
<tr>
<th>Social Context</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Definition</td>
<td>413465009 Afro-Caribbean (ethnic group)</td>
</tr>
<tr>
<td></td>
<td>116060000 Eating habit (life style)</td>
</tr>
<tr>
<td></td>
<td>24413000 Carpenter, general (occupation)</td>
</tr>
<tr>
<td></td>
<td>133932002 Caregiver (person)</td>
</tr>
<tr>
<td></td>
<td>415794004 Unknown racial group (racial group)</td>
</tr>
<tr>
<td></td>
<td>61154002 Hinduism (religion/philosophy)</td>
</tr>
<tr>
<td></td>
<td>22575004 Middle class economic status (social concept)</td>
</tr>
</tbody>
</table>
### Special Concept

**Definition**

<table>
<thead>
<tr>
<th>Inactive concept</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>363664003</td>
<td>Erroneous concept (inactive concept)</td>
</tr>
<tr>
<td>394899003</td>
<td>Oral administration of treatment (navigational concept)</td>
</tr>
</tbody>
</table>

**Navigational concept**

- 363743006 | Navigational concept (navigational concept) |

**Inactive concepts**

Inactive concepts are no longer active in the terminology. When the first release format (RF1) was used, inactivated content was moved into this hierarchy. However, this approach is no longer used in the current release format (RF2).

**Navigational concepts**

The concepts in navigational hierarchies are used for structured data entry and support the location of concepts in hierarchies. They can order data by priority or another convention (e.g. cranial nerve order or topics related to diabetes). Navigational concepts exist only to support navigation.

#### NO LONGER SUPPORTED IN SNOMED CT CORE

Navigational concepts:

- Are not suitable for recording or aggregating information
- Are direct subtypes of the concept 363743006 | Navigational concept (navigational concept) |
- Have no other supertype or subtype relationships
- Are linked to other concepts only by navigational links

For more information on navigational concepts, click here.

### Specimen

**Definition**

<table>
<thead>
<tr>
<th>Definition</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Entities that are obtained (usually from patients) for examination or analysis</td>
<td>384744003</td>
</tr>
<tr>
<td>122880004</td>
<td>Urine specimen obtained by clean catch procedure (specimen)</td>
</tr>
</tbody>
</table>

Specimen concepts can be defined by attributes which specify the:

- Normal or abnormal body structure from which they are obtained
- Procedure used to collect the specimen
- Source from which it was collected
- Substance of which it is comprised

**Specimen Attributes Summary**

When authoring in this domain, these are the approved attributes and allowable ranges. They are from the Human Readable Concept Model (HRCM).

#### Domain Information for 123038009 | Specimen (specimen) |

**Domain Constraint**

```plaintext
< 123038009 | Specimen (specimen) |
```

**Parent Domain**

-
### Specimen Defining Attributes

The following defining attributes correspond to the *Specimen Attributes Summary* table.

**Specimen source identity**

Specimen source identity specifies the type of individual, group, or physical location from which a specimen is collected.

For example,
- \[419695002\) |Environmental swab (specimen)\] has the Specimen source identity, Environment (environment)

**Specimen source morphology**

Specimen source morphology specifies the morphologic abnormality from which a specimen is obtained.

For example,
- \[447407009\) |Specimen from necrotic tissue (specimen)\] has the Specimen source morphology, Necrosis (morphologic abnormality)

**Specimen source topography**

Specimen source topography specifies the body site from which a specimen is obtained.

For example,
• 16209771000119101 | Specimen from left lower lobe of lung obtained by bronchoalveolar lavage procedure (specimen) | has the Specimen source topography, Segment of lower lobe of left lung (body structure)

Specimen procedure

Specimen procedure identifies the procedure by which a specimen is obtained.

For example,

• 384744003 | Lymph node from sentinel lymph node dissection and axillary dissection (specimen) | has the Specimen procedure, Dissection procedure (procedure)

Specimen substance

Specimen substance specifies the type of substance, pharmaceutical/biologic product, or physical object of which a specimen is comprised.

For example,

• 110897001 | Bone marrow cytological material (specimen) | has the Specimen substance, Bone marrow fluid (substance)

Staging and Scales

<table>
<thead>
<tr>
<th>Definition</th>
<th>Examples</th>
</tr>
</thead>
</table>
| This hierarchy contains concepts which are named, authoritative, and internationally relevant staging or grading systems used to either make a judgment about the patient, e.g. cognition, or, evaluate a patient to determine the phase, or progression of a disease. | Assessment  
- 273472005 | Functional status index (assessment scale)  
Staging  
- 254294008 | Tumor-node-metastasis (TNM) head and neck tumor staging (tumor staging) |

Some diseases are represented using a staging and/or grading system to signify the severity, extent, or rate of growth of a disease. For example, chronic kidney disease is represented with five stages determined by level of kidney function.

Assessment scale requests

- Generally, requests to add the most recent version of an assessment scale are accepted.
- Updated versions of existing content are also accepted.
- Older versions may be added if justification is appropriate. Older versions may also remain as active concepts due to the need to retain history on the use of specific instruments.

Modeling

Concepts of the type | Assessment using X assessment scale | are modeled with a proximal primitive parent of 445536008 | Assessment using assessment scale (procedure) or one of its subtypes, as appropriate.

For example,

- 445719003 | Assessment using visual analog pain scale (procedure) | has a parent of 445536008 | Assessment using assessment scale (procedure)

(See also Why is Content Rejected page, Proprietary Names for information about use of Questionnaire and Scale names)
Substance

<table>
<thead>
<tr>
<th>Definition</th>
<th>Examples</th>
</tr>
</thead>
<tbody>
<tr>
<td>Active chemical constituents of allergens, agents, substances, chemicals,</td>
<td>• 89889006</td>
</tr>
<tr>
<td>drugs, and materials (not Pharmaceutical/Biological Products)</td>
<td>• 52454007</td>
</tr>
</tbody>
</table>

Concepts from the Substance hierarchy are used to represent general substances and chemical constituents of Pharmaceutical / biologic products, which are in a separate hierarchy.

Substance concept and causative agent

When creating a new concept that includes a substance in the FSN, where no exact matching substance concept exists, then a new substance concept with an FSN, and a PT matching the FSN, should be created (the terms in the new concept should match the terms used in the FSN and PT of the substance concept selected as the causative agent).

For example,

- 418689008 |Allergy to grass pollen (disorder)| modeled with causative agent of 256277009 |Grass pollen (substance)|

Substance Attribute Summary

When authoring in this domain, this is the approved attribute and allowable range. It is from the Human Readable Concept Model (HRCM).

**Domain Information for 105590001 |Substance (substance)|**

| Domain Constraint | << 105590001 |Substance (substance)| |
| Parent Domain     | -            |
| Proximal Primitive Constraint | << 105590001 |Substance (substance)| |
| Proximal Primitive Refinement | - |

**Author View of Attributes and Ranges for 105590001 |Substance (substance)|**

<table>
<thead>
<tr>
<th>Attribute</th>
<th>Grouped</th>
<th>Cardinality</th>
<th>In Group Cardinality</th>
<th>Range Constraint</th>
</tr>
</thead>
<tbody>
<tr>
<td>726542003</td>
<td>Has disposition (attribute)</td>
<td>0</td>
<td>0..*</td>
<td>0..0</td>
</tr>
<tr>
<td>738774007</td>
<td>Is modification of (attribute)</td>
<td>0</td>
<td>0..*</td>
<td>0..0</td>
</tr>
</tbody>
</table>

Substance Defining Attributes

The following defining attributes correspond to the Substance Attribute Summary table.
Is modification of
This attribute indicates that the concept is a structural modification of another concept.

Has disposition
This attribute enables the creation of an association between a substance concept and a disposition (A disposition is defined as a behavior that a substance will exhibit or participate in, given the appropriate context)

Has disposition Overview
A new hierarchy, 726711005 |Disposition (disposition)|, was created to support the remodeling of the Substance hierarchy. The concepts in this hierarchy are used as the attribute value for the |Has disposition (attribute)| for concepts in the Substance hierarchy, to sufficiently define grouper concepts representing dispositions, and to sufficiently define concepts in other SNOMED CT hierarchies. To provide adequate context to differentiate concepts in this hierarchy from similar concepts in other existing SNOMED CT hierarchies, a new (disposition) semantic tag was created.

Modeling (stated view)

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>726711005</th>
<th>Disposition (&lt;&lt;)</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Exceptions: none</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(disposition)</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Definition status</th>
<th>900000000000074008</th>
<th>Necessary but not sufficient concept definition status (core metadata concept)</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Exceptions: none</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attributes</th>
<th>Not applicable</th>
</tr>
</thead>
</table>

Terming Guidelines
General Terming Guidelines
The following words should be avoided unless specifically identified as an exception in the editorial guidelines.

- agent
- analog
- and
- and/or
- content(s)
- derivative
- material
- metabolite
- modification
- or
- preparation
- product
- substance

Exceptions:
- Alkylating agent (disposition)
- Chelating agent (disposition)

Descriptions should be singular, not plural.

Note: Additional examples of types of dispositions will be added as needed.
Note: Additional examples of types of dispositions will be added as needed.

<table>
<thead>
<tr>
<th>Disposition representing:</th>
<th>FSN</th>
<th>Patterns:</th>
</tr>
</thead>
<tbody>
<tr>
<td>• receptor agonist or partial agonist</td>
<td></td>
<td>• X receptor agonist (disposition)</td>
</tr>
<tr>
<td>• receptor antagonist or partial antagonist</td>
<td></td>
<td>• X receptor antagonist (disposition)</td>
</tr>
<tr>
<td>• enzyme inhibitor or enzyme system inhibitor</td>
<td></td>
<td>• X inhibitor (disposition)</td>
</tr>
<tr>
<td>• enzyme activator</td>
<td></td>
<td>• X activator (disposition)</td>
</tr>
</tbody>
</table>

Example of receptor agonist:

- Dopamine receptor agonist (disposition)
- Opioid receptor agonist (disposition)
- Opioid receptor partial agonist (disposition)

Example of receptor antagonist:

- Beta-adrenergic receptor antagonist (disposition)
- Histamine receptor antagonist (disposition)
- Opioid receptor antagonist (disposition)
- Opioid receptor partial antagonist (disposition)

Example of inhibitor (enzyme):

- Acetylcholinesterase inhibitor (disposition)
- Phosphodiesterase 5 inhibitor (disposition)

Example of inhibitor (enzyme system):

- Hydrogen/potassium adenosine triphosphatase enzyme system inhibitor (disposition)
- Selective serotonin reuptake inhibitor (disposition)

Example of activator:

- Plasminogen activator (disposition)
### Preferred Term Patterns:

- $X$ receptor agonist
- $X$ receptor antagonist
- $X$ inhibitor
- $X$ activator

**Example of receptor agonist:**

- Dopamine receptor agonist
- Opioid receptor agonist

**Example of receptor antagonist:**

- Beta-adrenergic receptor antagonist
- Histamine receptor antagonist
- Opioid receptor antagonist

**Example of enzyme inhibitor:**

- Acetylcholinesterase inhibitor
- Phosphodiesterase 5 inhibitor
- Centrally acting acetylcholinesterase inhibitor

**Example of enzyme system inhibitor:**

- Proton pump inhibitor
- Selective serotonin reuptake inhibitor

**Example of enzyme inducer:**

- Plasminogen activator

**Exceptions:**

- Preferred terms may reflect the common clinical description for a disposition.
  
  **Example:**
  
  - [Hydrogen/potassium adenosine triphosphatase enzyme system inhibitor (disposition)] has Preferred Term [Proton pump inhibitor]
  - [3-Hydroxy-3-methylglutaryl-coenzyme A reductase inhibitor (disposition)] has Preferred Term [HMG-CoA reductase inhibitor]

**Synonyms**

- Synonyms are not allowed unless explicitly identified as an exception.
  
  **Exceptions:** None

### Disposition representing:

- response induced in an organism by a substance

**FSN Pattern:**

- $X$ (disposition)

**Example:**

- Growth factor (disposition)
- Cytokine (disposition)
- Carcinogen (disposition)
- Potassium channel blocker (disposition)
- Calcium channel blocker (disposition)
- Fusion inhibitor (disposition)
<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• X</td>
</tr>
</tbody>
</table>

Example:
- Growth factor
- Cytokine
- Carcinogen
- Potassium channel blocker
- Calcium channel blocker
- Fusion inhibitor

<table>
<thead>
<tr>
<th>Synonyms</th>
<th>Synonyms are not allowed unless explicitly identified as an exception.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Exceptions: none identified</td>
</tr>
</tbody>
</table>

**Disposition representing:**
- a chemical effect or physical property

<table>
<thead>
<tr>
<th>FSN Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td>• X (disposition)</td>
</tr>
</tbody>
</table>

Example:
- Alkylating agent (disposition)
- Chelating agent (disposition)

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• X</td>
</tr>
</tbody>
</table>

Example:
- Alkylating agent
- Chelating agent

<table>
<thead>
<tr>
<th>Synonyms</th>
<th>Synonyms are not allowed unless explicitly identified as an exception.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Exceptions: none identified</td>
</tr>
</tbody>
</table>

**Exemplar**

The following illustrates the **stated** and **inferred** view:
## Substance Naming and Modeling Conventions

### General Assumptions and Scope for Substance Hierarchy

**General Assumptions and Requirements**

General assumptions and requirements include the following:

<table>
<thead>
<tr>
<th>Assumption or Requirement</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>1</strong></td>
</tr>
</tbody>
</table>
| The |Substance| hierarchy contains concepts that can be used for recording chemical constituents of drug products, food and chemical allergens, adverse reactions, toxicity or poisoning information, and physicians and nursing orders. Concepts from this hierarchy represent general substances and chemical constituents of |Pharmaceutical / biologic product (product)| which are in a separate hierarchy. However, sub-hierarchies of |Substance| also include but are not limited to: |Body substance (substance)| (concepts to represent body substances) and |Diagnostic substance (substance)|.  

**Examples of Substance concepts:**  
- Insulin (substance)  
- Methane (substance)  
- Chromatin (substance)  
- Albumin (substance)  
- Endorphin (substance)  
- Paracetamol (substance) |
| **2**                     |
| Concept model conforms to description logic principles, including use of the classifier to organize the concepts in the hierarchy where appropriate  
  - Top level concepts in the hierarchy will primarily be grouper concepts for both the stated and inferred view |
| **3**                     |
| Concept model includes attributes necessary to define concepts where appropriate to ensure consistent and reproducible modeling of concepts |
| **4**                     |
| Concepts representing dispositions in the |Substance| hierarchy shall be sufficiently defined using proximal primitive modeling methodology unless explicitly noted as an exception in the editorial guidelines  
  - Additional information about proximal primitive modeling can be found in the SNOMED CT Editorial Guide  
  - Inclusion of intermediate primitives or manually maintained IS A relationships will negatively impact the ability to maintain the content and to identify equivalencies; further information can be found in the SNOMED CT Editorial Guide. |

**Scope of Content**

This section applies to the |Substance| hierarchy in the International Release:

<table>
<thead>
<tr>
<th>In Scope</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>1</strong></td>
</tr>
<tr>
<td>Substances used in medicinal products</td>
</tr>
<tr>
<td><strong>2</strong></td>
</tr>
<tr>
<td>Substances that support the modeling of concepts in other SNOMED CT hierarchies</td>
</tr>
</tbody>
</table>
**In Scope**

3 Substances that support the modeling of the existing herbal products in SNOMED CT  

Note: 418165002 |Herbal medicine agent (substance)| is a role and will be replaced in future. Its descendants will be retained “as is” until use cases and/or detailed requirements are known. Requests for addition of new concepts or for modification of existing concepts will be evaluated on a case-by-case basis.

**Out of Scope**

1 Concepts that refer to dose form (e.g. solution) or route of administration (e.g. topical)  
   - Existing instances will be inactivated as nonconformant to editorial policy. Requests for new instances will be rejected

2 Concepts that refer to role or a specific context (e.g. dietary, medicinal, non-pharmaceutical, substance of abuse)  
   - Existing instances will be inactivated as nonconformant to editorial policy. Requests for new instances will be rejected

3 Concepts that refer to a release state (e.g. immediate release, extended release)  
   - Existing instances will be inactivated as nonconformant to editorial policy. Requests for new instances will be rejected

4 Concepts that refer to a brand or trade name  
   - Existing instances will be inactivated as nonconformant to editorial policy. Requests for new instances will be rejected

5 Concepts that state "total" due to the inability to differentiate the definition between the "base" substance and "total" substance (e.g. "cholesterol" versus "total cholesterol") as well as the inability to create an appropriate relationship between the "base" and "total" concepts  
   - Existing instances will be inactivated as nonconformant to editorial policy. Requests for new instances will be rejected

6 Substances of the pattern |X molecule of Y organism (substance)| which reference an organism that is not of interest to human medicine are considered out of scope of SNOMED CT.  
   - Example: |Ribonucleic acid of Porcine reproductive and respiratory syndrome virus (substance)|  
   - Existing instances will be inactivated as nonconformant to editorial policy. Requests for new instances will be rejected

7 Substances used in homeopathic products  
   - Existing instances will be retained for now; however requests for new instances will be rejected

8 Substances used in traditional medicine products  
   - Existing instances will be retained for now; however requests for new instances will be rejected

Note: Content requests related to the areas where terming and modelling guidelines are required (e.g. vaccine) will be accepted and placed in Inception/Elaboration status until a long term plan is agreed.
Out of Scope

<p>| | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>9</td>
<td>Concepts that represent a combination of two or more separate substances</td>
</tr>
<tr>
<td></td>
<td>- Existing instances have been inactivated. Changes to SNOMED CT concept model to permit the use of concepts within the products hierarchy (e.g. vaccine products) as well as use of General Concept Inclusion functionality to model disorder and findings concepts means that these combined substances concepts will no longer be required. Requests for new instances will be rejected.</td>
</tr>
</tbody>
</table>

Groupers for Substance Hierarchy

Substance hierarchy grouper concepts
In the SNOMED CT substance hierarchy there are a large number of grouper concepts. Currently groupers within the substances can be considered to be role based, dispositions or structural groupers.

The intent is to build the SNOMED CT substances hierarchy along structural or disposition groupers and use a HAS_DISPOSITION relationship to define grouper concepts in the substances hierarchy. Grouper concepts that do not reference structural or disposition/functional properties of a substance should not be included within the substances hierarchy and existing role concepts are considered for relocation outside of the substances hierarchy.

Structure based Groupers
Groupers that organize substances by their chemical structure are used as the main hierarchy for substances. Where substances are heterogeneous and do not have a single identifiable chemical structure such as those of biological origin it may be more appropriate to organize them by source.

Dispositions Groupers
In the context of substances, a disposition is “a behavior that a substance will exhibit or participate in, given the appropriate context.” This context-based definition of disposition would allow us to assign HAS-DISPOSITION values that are necessarily true, even though the substance does not exhibit the disposition in all contexts.

Role based Groupers
Role based groupers are associated with a particular purpose or outcome. Roles are a function of the way the substance is formulated or presented and so may not be applicable to all products containing that substance. For this reason role based groupers should not be included in the substances hierarchy.

Restructure of the top level substances hierarchy
Previously, the substances hierarchy in SNOMED CT was organized using a number of different axes some of which did not comply with the terminological principle of being always true. This resulted in incorrect inferences both within the substance hierarchy and other hierarchies where substances were used to define concepts. To avoid this, the substances hierarchy has been organized by characteristics that are always true (e.g. chemical structure). For substances where the structure is unknown or the substance is heterogeneous, other characteristics that are always true should be used as parent concepts within the substances hierarchy.

The current substances top level hierarchy includes a number of grouper concepts that are role or use case based. The proposal is to create a substances top level hierarchy as below.
The grouper concepts that are retained as top level substance concepts will need review of their descendants, but ultimately it should provide a consistent hierarchy.

There are similarities in the scope of "ISO 11238 Health informatics – Identification of medicinal products – Data Elements and Structures for the Unique Identification and Exchange of Regulated Information on Substances" and the SNOMED CT substances revision, since both consider a substance to be defined by properties such as molecular structure and not by how it is formulated or used.

IDMP groups substances as being one of five types of single substances:

- (simple) chemical
- protein/peptide
- nucleic acid
- polymer
- structurally-diverse

or they are classified as a mixture. In IDMP the definition of a mixture is much broader that would be perceived by clinical users. IDMP defines starches as mixtures and also compounds such as isophane insulin and gentamicin sulfate as mixtures. This broad definition for mixtures would mean that many of the SNOMED CT substances would be classified as mixtures. These IDMP types are not represented in SNOMED CT since they are not mutually exclusive and are open to some interpretation. The IDMP documentation does provide guidance on the selection of the correct type for a substance but also acknowledges that a single substance may have two separate type definitions. In addition, the use of the word “mixture” in the substances hierarchy is to be discouraged, since as a general scope statement SNOMED CT does not include combination substances that are a mixture of more than one individual chemical even when the two exhibit a synergistic effect. E.g. tazobactam and piperacillin.

Changes to SNOMED CT concept model in the future to permit the use of concepts within the products hierarchy to model disorder and findings concepts means that these combined substances concepts will no longer be required and so will be inactivated at that time.

Differences between SNOMED CT substances and IDMP Substances

1. The scope of substances in SNOMED CT is broader than that of IDMP, since the definition of regulated products is not the only use case supported by the SNOMED CT substances hierarchy.
2. IDMP classes are used to identify which features could/should be identified as defining. This is different from the SNOMED CT requirement to identify top level concepts to support a hierarchy that provides a logical structure for substance concepts.
3. IDMP makes a distinction between Substance and the more closely defined Specified Substance. Substances are "any matter of defined composition that has discrete existence, whose origin may be biological, mineral or chemical".
   a. Specified Substances are "defined by groups of elements that describe multi-substance materials or specifies further information on substances relevant to the description of Medicinal Products".
   b. Specified substances include mixture substances, substances defined by pharmacopoeial specification or substances where a particular manufacturing process is specified.
In order for SNOMED CT to support the IDMP work, whose primary goal is to define unambiguously all substances present in regulated products, it is necessary for both the concepts that are IDMP Substances and those that are IDMP Specified Substances to be present in the SNOMED CT Substance hierarchy. Stereoisomers, hydrates and solvates will be included in the SNOMED CT substances hierarchy as concepts as well as concepts to represent their base chemical in the SNOMED CT substance hierarchy. Since both IDMP Substances and IDMP Specified Substances are candidate concepts to be used in the ingredient role attributes it does not add value in the definition of these concepts to separately identify IDMP Substances and IDMP Specified Substances in SNOMED CT and would likely cause confusion where substance concepts are used to define concepts outside the scope of IDMP and the medicinal product hierarchy.

Substance Groupers Based on Structure

Overview

Groupers based on structural properties of the substance that are deemed to be clinically useful will be included in the Substance hierarchy.

Example:
- Benzodiazepine (substance)
- Quinolone (substance)

Modeling (stated view)

| Parent concept | Most distal appropriate descendant of 312413002|Substance categorized by structure (substance)|
|----------------|-------------------------------------------------|
|                | • Exceptions: none identified                    |
| Semantic tag   | (substance)                                      |
| Definition status | 9000000000000000074008|Necessary but not sufficient concept definition status (core metadata concept)|
|                | • Exceptions: none identified                    |
| Attributes     | None                                            |
|                | • Exceptions: none                              |

Exemplar for Concept Model

The following illustrates the stated and inferred view.
Terming Guidelines

<table>
<thead>
<tr>
<th>FSN</th>
<th>Use the following pattern for the FSN. Groupers should be expressed in the singular. (Compound not compounds)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Example for substances that represent a structural grouper:</td>
</tr>
<tr>
<td></td>
<td>- Organic nitrogen compound (substance)</td>
</tr>
<tr>
<td></td>
<td>- Halide compound (substance)</td>
</tr>
<tr>
<td></td>
<td>- Triazine derivative (substance)</td>
</tr>
<tr>
<td></td>
<td>- Copper and/or copper compound (substance)</td>
</tr>
<tr>
<td></td>
<td>- Phenothiazine and/or phenothiazine derivative (substance)</td>
</tr>
<tr>
<td></td>
<td>- Nucleotide (substance)</td>
</tr>
<tr>
<td>Where the same word is used to identify both a group of substances</td>
<td>Substances with X structure (substance)</td>
</tr>
<tr>
<td>with a particular chemical structure and a particular chemical</td>
<td>Examples:</td>
</tr>
<tr>
<td>substance the following terming for the FSN will be used</td>
<td>- Substance with tetracycline structure (substance)</td>
</tr>
<tr>
<td>for the substance grouper concept.</td>
<td>- Substance with cresol structure (substance)</td>
</tr>
<tr>
<td></td>
<td>- Substance with benzene structure (substance)</td>
</tr>
</tbody>
</table>
### Preferred Term

Use the following pattern for the Preferred Term,

Example:

- Organic nitrogen compound
- Halide compound
- Triazine derivative
- Copper and/or copper compound
- Phenothiazine and/or phenothioazine derivative
- Nucleotide

Where the same word is used to identify both a group of substances with a particular chemical structure and a particular chemical substance the following pattern will be used for PT the substance grouper concept.

Substance with X structure

Examples:

- Substance with tetracycline structure
- Substance with cresol structure
- Substance with benzene structure

### Synonyms

A synonym to match the FSN must be created.

Additional synonyms are not allowed unless explicitly identified as an exception in the Editorial Guidelines.

---

### Exemplar for Hierarchy

**Parents**

- Nitrogen compound (substance)
- Organic compound (substance)

```plaintext
<table>
<thead>
<tr>
<th>Organic nitrogen compound (substance)</th>
</tr>
</thead>
<tbody>
<tr>
<td>SCTID: 299979007</td>
</tr>
<tr>
<td>299979007</td>
</tr>
<tr>
<td>en Organic nitrogen compound (substance)</td>
</tr>
</tbody>
</table>
```

- Azo derivative (substance)
- Amaranth (substance)
- Azo dye (substance)
- Azodicarbonamide (substance)
- Azodilsobutyrodinitrile (substance)
Figure 1: Browser view of 299979007 | Organic nitrogen compound (substance) |

**Parents**
- Heavy metal and/or heavy metal compound (substance)

**Children (6)**
- Copper (substance)
- Copper compound (substance)
- Copper dust and mist (substance)
- Copper fumes (substance)
- Copper isotope (substance)
- Copper radioisotope (substance)
- Free copper (substance)
Figure 2: Browser view of 767209009 | Copper and/or copper compound (substance) |

Parents
- Organic cyclic compound (substance)

Substance with cresol structure (substance)
SCTID: 766224008

766224008 | Substance with cresol structure (substance) |
  en Substance with cresol structure (substance)
  en Substance with cresol structure

Children (7)
- Amy/melacresol (substance)
- Chlorocresol (substance)
- Cresol (substance)
- Cyclofenil (substance)
- M-cresol (substance)
- Tetrabromo-o-cresol (substance)
- Tetrachloro-1,4- benzoquinone (substance)
Figure 3: Browser view of 766224008 |Substance with cresol structure (substance)|

Substance Groupers Based on Disposition
Overview
Groupers based on disposition that are deemed to be clinically useful and that can be sufficiently defined will be included in the |Substance| hierarchy. Disposition is defined as a behavior that a substance will exhibit or participate in, given the appropriate context.

There is no requirement to introduce a new semantic tag in order to distinguish concepts representing a substance disposition from any other type of concept in the |Substance| hierarchy.

Note: This section applies to concepts representing a single disposition. It does not apply to concepts representing a disposition combined with a structural grouper or concepts representing more than one disposition.

Modeling (stated view)

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>105590001</th>
<th>Substance (substance)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Exceptions: while the goal is to model these concepts using proximal primitive modeling, there will be a transition over several release cycles to get to that state. In the meantime, concepts may have stated parents other than 105590001</td>
<td>Substance (substance)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Definition status</th>
<th>900000000000073002</th>
<th>Sufficiently defined concept definition status (core metadata concept)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Exceptions: none</td>
<td></td>
</tr>
<tr>
<td>Has disposition</td>
<td></td>
<td></td>
</tr>
<tr>
<td>----------------</td>
<td></td>
<td></td>
</tr>
<tr>
<td>- Range: 726711005</td>
<td>Disposition (disposition) - descendants only</td>
<td></td>
</tr>
<tr>
<td>- NOTE: While the allowed range is broader, the Substance Groupers based on Disposition should only use descendants of the concept 726711005</td>
<td>Disposition (disposition) as the attribute value.</td>
<td></td>
</tr>
<tr>
<td>- Exceptions: none identified</td>
<td></td>
<td></td>
</tr>
<tr>
<td>- Cardinality: 1..1</td>
<td></td>
<td></td>
</tr>
<tr>
<td>- NOTE: While the allowed range is broader, the Substance Groupers based on Disposition should have one and only one</td>
<td>Has disposition</td>
<td>attribute.</td>
</tr>
<tr>
<td>- Exceptions: none identified</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Note: Substance groupers representing etiopathic dispositions (e.g. 88376000 | Carcinogen (substance), 45986006 | Teratogen (substance)) will be created. Since most substances can exhibit these characteristics in some circumstances, the descendant concepts will not be populated. Exceptions will be noted in the editorial guidelines.

Exemplar for Grouper Concept Based on Disposition
The following illustrates the **stated** view for grouper concepts based on disposition.

![Diagram](image1)

The following illustrates the **inferred** view for grouper concepts based on disposition.

![Diagram](image2)

Terming Guidelines

| FSN | Use the following pattern for the FSN. The FSN must reflect the terming used to describe the disposition consistent with the terming of the | Has disposition | attribute value. |
|-----|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
|     | Example for dispositions that represent a mechanism of action: |
|     | - Substance with opioid receptor agonist mechanism of action (substance) |
|     | - Substance with histamine receptor antagonist mechanism of action (substance) |
|     | - Substance with acetylcholinesterase inhibitor mechanism of action (substance) |
Preferred Term

Use the following pattern for the Preferred Term. The Preferred Term must reflect the terming used to describe the disposition consistent with the terming of the |Has disposition| attribute value.

Example for dispositions that represent a mechanism of action:

- Opioid receptor agonist
- Histamine receptor antagonist
- Acetylcholinesterase inhibitor

Synonyms

Synonyms corresponding to the FSN are not required.

Additional synonyms are allowed only if they are consistent with the synonyms for the corresponding disposition concept.

Substance Groupers Based on Both Structure and Disposition

Overview

Groupers based on both structure and disposition that are deemed to be clinically useful and that can be sufficiently defined will be included in the |Substance| hierarchy.

Note: This section applies to concepts representing a single structural parent and a single disposition. It does not apply concepts representing more than one structural parent or disposition.

Modeling (stated view)

<table>
<thead>
<tr>
<th>Stated parent concept</th>
<th>The stated parent concept must be the concept that represents the structural grouper. If such a concept does not exist, it must be created and its stated substance descendants added before the grouper based on both structure and disposition can be created.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Exceptions: none</td>
</tr>
<tr>
<td>Definition status</td>
<td>900000000000073002</td>
</tr>
<tr>
<td></td>
<td>• Exceptions: none</td>
</tr>
<tr>
<td>Attribute:</td>
<td>Range: 726711005</td>
</tr>
<tr>
<td>Has disposition</td>
<td>• NOTE: While the allowed range is broader, the Substance Groupers based on both Structure and Disposition should only use descendants of the concept 726711005</td>
</tr>
<tr>
<td></td>
<td>• Exceptions: none identified</td>
</tr>
<tr>
<td></td>
<td>• Cardinality: 1..1</td>
</tr>
<tr>
<td></td>
<td>• Exceptions: none identified</td>
</tr>
</tbody>
</table>

Exemplar for Grouper Concept Based on both Structure and Disposition

The following illustrates the stated and inferred view for grouper concepts based on both structure and disposition.
Figure 1: Stated view of 438942000 | Piperidine derivative with histamine receptor antagonist mechanism of action (substance)|
### Terming Guidelines

<table>
<thead>
<tr>
<th><strong>FSN</strong></th>
<th>Use the following pattern for the FSN, with X representing the structure and Y representing the disposition. The FSN should align with the FSN for the substance grouper and disposition used as the stated parent and attribute value respectively.</th>
</tr>
</thead>
<tbody>
<tr>
<td>X with Y mechanism of action (substance)</td>
<td>Example:</td>
</tr>
<tr>
<td>- Piperazine derivative with histamine H1 receptor antagonist mechanism of action (substance)</td>
<td></td>
</tr>
<tr>
<td>- Substance with dihydropyridine derivative structure and calcium channel blocker mechanism of action (substance)</td>
<td></td>
</tr>
<tr>
<td>- Substance with organophosphorus structure and acetylcholinesterase inhibitor mechanism of action (substance)</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>Preferred Term</strong></th>
<th>Use the following pattern for the Preferred Term, with X representing the structure and Y representing the disposition. The FSN should align with the FSN for the substance grouper and disposition used as the stated parent and attribute value respectively.</th>
</tr>
</thead>
<tbody>
<tr>
<td>X Y</td>
<td>Example:</td>
</tr>
<tr>
<td>- Piperazine derivative histamine H1 receptor antagonist</td>
<td></td>
</tr>
<tr>
<td>- Dihydropyridine derivative calcium channel blocker</td>
<td></td>
</tr>
<tr>
<td>- Organophosphorus acetylcholinesterase inhibitor</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>Synonyms</strong></th>
<th>A synonym to match the FSN is not required. Additional synonyms are allowed only if they are consistent with the synonyms for the corresponding structure grouper and disposition.</th>
</tr>
</thead>
</table>

### Concepts Representing a Substance or its Modifications

**Overview**
The substance redesign project has made a number of decisions regarding the representation of a substance and its modifications.

1. Modification concepts that exist to group modifications or derivatives of a specific substance not be created.
2. Metabolite groupers in the substances hierarchy are considered roles. No further concepts of this format will be created in the substances hierarchy. Existing concepts will only be retained where there is a specific requirement and will be modeled as a child of the concept Metabolite (substance). See relative section elsewhere in this document.
3. Structural groupers that reference modifications as a chemical group will be retained. Where the structural group name also refers to a specific chemical the grouper will be re-termed “Substance with X structure”. See relative section elsewhere in this document.
4. There is no requirement to introduce a new semantic tag in order to distinguish concepts representing a substance or its modifications from any other type of concept in the |Substance| hierarchy.

### Modeling using the IS Modification attribute (stated view)

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(substance)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Definition status</td>
<td>900000000000074008 [Necessary but not sufficient concept definition status (core metadata concept)]</td>
</tr>
<tr>
<td></td>
<td>• Exceptions: none identified</td>
</tr>
<tr>
<td>Attributes</td>
<td>None</td>
</tr>
<tr>
<td></td>
<td>• Exceptions: none</td>
</tr>
</tbody>
</table>

### Exemplar

The following illustrates the **stated** view

The following illustrates the **inferred** view for concepts that are a modification.
Guidelines for the use of the IS Modification attribute with Examples

Substances may have zero to many IS modification attribute(s)

For example, Morphine hydrochloride (substance)

For example, Fluorothymidine (18-F) (substance)
For example, Doxepin hydrochloride (substance)

Hydrates have a IS MODIFICATION relationship to the unspecified salt

For example, Caffeine hydrate (substance) IS MODIFICATION of Caffeine (substance)

This is the described view.
This is the inferred view

For example, Clindamycin hydrochloride monohydrate (substance) IS MODIFICATION of Clindamycin hydrochloride (substance)

This is the stated view:

This is the inferred view:
For example, Calcium lactate pentahydrate (substance) IS MODIFICATION of Calcium lactate (substance)
This is the **stated and inferred** view.

Anhydrous salts have a IS MODIFICATION relationship to the unspecified salt
For example, Theophylline anhydrous (substance) IS MODIFICATION of Theophylline (substance)
This is the **stated** view
This is the **inferred** view
The salts are modeled with an IS Modification to the acid substance.

For example, Pamidronate monosodium (substance) IS MODIFICATION of Pamidronic acid (substance). This is the stated and inferred view.

Liposomal preparations are modifications of the chemical substance.

For example, Daunorubicin citrate liposome (substance).
Pegylated substances should be modelled as "Is_modification" of generic substance—if such a substance has been made available; not all pegylated medicinal substances have a non-pegylated form as the non-pegylated form may have been too immunogenic or too toxic.

For example, 785674001 |Turoctocog alfa pegol (substance)|

Concepts specifying a particular physical form (e.g. micronized, macrocrystal, microsphere) should have a parent concept that relates to the structure of the substance and also an Is modification of (attribute) relationship to the unspecified substance concept.
There should be an “is_modification” relationship between the substance and its esterified form.
Grouper concepts should not be targets of an IS Modification relationship.

There is no IS MODIFICATION relationship between stereoisomers. Each stereoisomer should be treated as a separate substance. E.g. Cetirizine and levocetirizine may be sibling concepts but do not have an IS MODIFICATION attribute relationship associating them.

There is no IS MODIFICATION relationship between "antibody-drug conjugate" (e.g. trastuzumab) as they are related to the delivery mechanism rather than modification of base concept.

"Is modification" relationship is not applicable to prodrugs. For example, aspirin is not generally considered as a modification of the substance salicylic acid. This is because there is no requirement for the "has active ingredient" attribute of medicinal products containing prodrug substances to use any relationship to the active substance to manage relationships within the medicinal product hierarchy.

"Is modification" relationship is not applicable between a glycan and its glycoconjugate
Exemplar: 259289005 [Trimethylene glycol (substance)] will not be a modification of 52086008 [Glycol (substance)].

Naming and Modeling Guidelines for Substance Hierarchy

This section of the document provides both general terming and modeling guidance for substance concepts and also guidance for specific types of substances.

Substance Concept General Guidelines

Overview

Modeling (stated view)

| Parent concept | <<105590001 | Substance (substance)| |
|----------------|-----------------|
| Exceptions: none |

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(substance)</th>
</tr>
</thead>
</table>
### Definition status

<table>
<thead>
<tr>
<th>Definition status</th>
<th>900000000000074008</th>
<th>Necessary but not sufficient concept definition status (core metadata concept)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>• Exceptions: none identified</td>
</tr>
</tbody>
</table>

### Attribute:

<table>
<thead>
<tr>
<th>Attribute</th>
<th>738774007</th>
<th>Is Modification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Is Modification</td>
<td>738774007</td>
<td>Is Modification</td>
</tr>
<tr>
<td>Range &lt;105590001</td>
<td>Substance (substance)</td>
<td></td>
</tr>
<tr>
<td>Cardinality: 0..*</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Exceptions: none identified</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Attribute:

<table>
<thead>
<tr>
<th>Attribute</th>
<th>726711005</th>
<th>Disposition (disposition)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Has disposition</td>
<td>726711005</td>
<td>Disposition (disposition)</td>
</tr>
<tr>
<td>Range: 726711005</td>
<td>Disposition (disposition)</td>
<td></td>
</tr>
<tr>
<td>- descendants only</td>
<td></td>
<td></td>
</tr>
<tr>
<td>NOTE: While the allowed range is broader, substance concepts should only use descendants of the concept 726711005</td>
<td>Disposition (disposition)</td>
<td></td>
</tr>
<tr>
<td>- as the attribute value.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardinality: 0..*</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Exceptions: none identified</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Naming Guidelines

#### General Naming Guidelines

Descriptions should be singular, not plural.

- Exception: Fumes should be expressed in the plural (i.e. fumes as opposed to fume).

Descriptions should not include additional descriptors that reference a use case.

For example,

- **Non-pharmaceutical**

Some regulatory agencies may use prefixes or suffixes to distinguish between different manufacturers of a single substance but these will not be used in SNOMED CT; the INN name will be used.

Example:

- The FDA uses "ado-trastuzumab"; the INN for this substance is "trastuzumab". The INN should be reflected as the US and GB names in SNOMED CT International Release because the prefix represents a specific manufacturer’s product.
- The FDA uses "emicizumab-kxwh" and "vestronidase alfa-vjbk"; the INN for these substances are "emicizumab" and "vestronidase alfa". The INN should be reflected as the US and GB names in SNOMED CT International Release because the suffix represents a specific manufacturer’s product.

Substance concepts should not include a strength.

Substance concepts should not include a dose form or use case

The FSN is aligned with the INN, the Preferred Terms are aligned with USAN and BAN (BAN almost always aligns with the INN). The Preferred Term in the US dialect variation must be given the acceptability of Acceptable (A) in the GB dialect and vice versa.

For isomers the INN names use the expanded prefix e.g. levo or dextro and so should be used for the FSN and Preferred Term.

Changes to descriptions allocated to existing substance concepts may impact on the terming of medicinal product concepts and so should be undertaken with caution.
General Naming Guidelines

For case sensitivity assignment:

- Greek alphabetical terms in substance concepts have been made case insensitive
- For substance terms that have a single letter - either a single lower case letter or a single upper case letter - the case sensitivity should reflect this either CS if the single capital letter or single lower case letter is at the beginning of the term, or CI if the single capital letter or single lower case letter is contained within an otherwise case insensitive term
- Proper names in substance terms should begin with a capital letter
- Numeric numbers in substance terms should be ignored for case significance. Simply, they won’t be displayed differently if we switch between upper and lower case. The case sensitivity decision should be based on the rest of the letters and words in the term. When modeling terms the word following the numbers does not begin with a capital letter e.g. correct terming is 5-hydroxytryptamine (ci) and not 5-Hydroxytryptamine, and the substance abbreviated to ‘6-TG’ is CS not CI.

Antibodies and antigens
Overview
This section includes concepts that represent Antigen of X organism, Antibody to X organism and Immunoglobulin G, M, A, E, D antibody to X organism.

- When referring to an organism name, while the Linnean taxon ranks (such as "genus") are not included, the subspecies variants (such as "biotype" and "serotype") are included in the naming. This is to avoid ambiguity when the same number or letter is used to refer to different organism variants. For example, without mentioning the specific variant (serogroup vs. serotype) and the nomenclature system (Danish vs. American), "Streptococcus pneumoniae 48" can refer to the following:
  - Streptococcus pneumoniae Danish serotype 48 (which is equivalent to Streptococcus pneumoniae American serotype 82)
  - Streptococcus pneumoniae American serotype 48 (which is equivalent to Streptococcus pneumoniae Danish serotype 7B)
  - Streptococcus pneumoniae serogroup 48

Modeling - Antigen of X organism

| Parent concept | Most distal appropriate descendant of 116633006 | Microbial antigen (substance) |
|----------------|-----------------------------------------------|
|                | Exceptions: none                               |

Semantic tag (substance)

Definition status 900000000000074008 | Necessary but not sufficient concept definition status (core metadata concept) |

Attribute N/A

Naming Guidelines - Antigen of X organism
The antigen name is usually derived from the biological name of the organism as opposed to the “disease" or "infection" that the organism may cause and/or that the vaccine may provide immunization for.
Antigen variants:
There is a requirement for the inclusion of characteristic technologies that are essential for distinguishing between different antigen variants that are used in manufacturing vaccine products e.g live attenuated, inactivated, or subunit antigens. Antigen variants are evaluated for:

- References such as WHO, CDC, UpToDate, vaccine package inserts.
- Inclusion in the international release vs. national extensions
- Hierarchy they belong to: Substance or Product.
  - If they belong in the Substance hierarchy, the classification with respect to other antigens (legacy content that does not adhere to the following guideline will be updated in future releases)

Variants reviewed to date:
- “Antigen of X” is in scope for the international release and is modeled in the Substance hierarchy. It is a generic grouper concept and subsumes all instances of antigen variants related to Organism X.
  - X refers to a bacteria, virus, fungus, or parasite except when antigen refers to the organism “toxin/toxoid”, where it can only apply to a bacteria.
“Antigen of live attenuated X” is in scope for the international release and is modeled in the Substance hierarchy as a direct child of Antigen of X.

- It refers to attenuated whole cell bacteria or whole virus where the strains are made less virulent so infection is usually inapparent or very mild. It may be used in the creation of vaccine products for certain patient groups and hence is of clinical significance.
- The following subtypes are also in scope for the international release and are modeled in the Substance hierarchy as direct children of Antigen of live attenuated X. Note that in the following examples "human" and "bovine" refer to source organisms which are differentiated from the organism producing antigen i.e. Rotavirus.
  - "Live attenuated human X" e.g. Antigen of live attenuated human Rotavirus serotype G1P[8]
  - "Live attenuated human-bovine reassortant X" e.g Antigen of live attenuated human-bovine reassortant Rotavirus serotype G1

“Antigen of whole inactivated X” is in scope for the international release and is modeled in the Substance hierarchy as a direct child of Antigen of X.

- It refers to the killed version of the organism that causes a disease.
- As of the July 2020 release, the word “whole” is added to all new and existing concepts referring to "Antigen of inactivated X" to further clarify the differentiation between "Antigen of whole inactivated organism" (referring to a killed organism as a whole) and "Antigen of organism [subunit]" (referring to subparts of an organism).

“Antigen of acellular X” is in scope for the international release and is modeled in the Substance hierarchy as the direct child of "Antigen of X".

- "Antigen of acellular X", when it exists, can only apply to a bacteria.

“Antigen of X [subunit]” is in scope for the international release and is modeled in the Substance hierarchy as the direct child of “Antigen of X”.

- Subunit antigens differ from inactivated whole-cell antigens by referring only to the antigenic parts of the pathogen. These parts are necessary to elicit a protective immune response. The word “inactivated” does not need to be part of the name unless it is referring to a whole cell or a whole virus i.e. for descriptions referring to a subunit, the word “inactivated” will be omitted as redundant.
- [Subunit] refers to a sub-part of an organism i.e. a specific, isolated protein of the pathogen, a recombinant protein (made by recombinant DNA techniques), an inactivated toxin (toxoid), or a capsular polysaccharide/oligosaccharide coating of an encapsulated bacterium:
  - "Antigen of X [protein/recombinant protein]" is in scope for the international release and is modeled in the Substance hierarchy.
    - It represents a specific, isolated protein of the pathogen or a recombinant protein.
    - When both "Antigen of X [protein]" and "Antigen of X [recombinant protein]" exist for X organism, they are modelled as siblings.
  - "Antigen of X capsular polysaccharide/oligosaccharide" is in scope for the international release and is modeled in the Substance hierarchy.
    - It represents a polysaccharide/oligosaccharide antigen and acts as a grouper for the following:
      - "Antigen of X capsular polysaccharide/oligosaccharide unconjugated", which represents a polysaccharide/oligosaccharide antigen without conjugation to any carrier protein or toxoid
      - "Antigen of X capsular polysaccharide/oligosaccharide conjugated", represents a grouper concept and will be created in the international release and in the Substance hierarchy per request and (based on current ED guidelines) only if it has more than one child concept.
    - Exception can apply if there is a use case that support retaining/creating these conjugate groupers:
      - Recording the history of a conjugated vaccine when the type of conjugated protein is not known
      - Aggregating data for forecasting
    - The existing grouper concepts with just one child will not be deprecated at this time. However, requests for addition of these concepts will be rejected.
• “Antigen of X capsular polysaccharide/oligosaccharide conjugated to Y”, which represents an oligosaccharide or polysaccharide antigen attached to a protein Y, where “Y” refers to a carrier protein to increase efficacy and immunogenicity e.g. Corynebacterium diphtheriae cross-reacting material 197 protein.
  - It is the direct child of single parent "Antigen of X capsular polysaccharide/oligosaccharide conjugated". There are no clinical use-case that requires association to the conjugated part of antigen as an additional parent. This classification is in line with all other similar concepts (modifications such as pegylated substances).
• “Antigen of X toxoid” is in scope for the international release and is modeled in the Substance hierarchy.
  - Like other subunit antigens, “Antigen of X toxoid” is classified under "Antigen of X". While the toxin is not an intrinsic part of the organism, it is a product of the organism that would not exist in the absence of that organism. In other words, there is always a direct association between the substance and the source organism. There are other antigenic proteins that are generated by the organism (e.g. surface protein) and they are classified as children of “Antigen X organism”. The only difference, in comparison to toxoids, is that they are not being excreted by the organism.
  - Toxoid antigens are based on the toxin produced by certain bacteria (e.g. tetanus or diphtheria), which has been chemically processed so that it is still immunogenic. Once the toxin has been inactivated, it is called a toxoid.
  - A toxoid can be an antigen in its own right, or it can be conjugated to another antigen.
• When referring to Organism parts/subunits:
  - Referring to more than one subunit (e.g. combined protein such as Bordetella pertussis FIM 2 and FIM 3 antigen) would not be acceptable for the Substance hierarchy i.e. the combined protein needs to be modelled at the product level with more than one active ingredient.
  - Abbreviated organism subunit names such as "Corynebacterium diphtheriae CRM197 protein", are not allowed in a fully specified name (and similar synonym). The FSN and similar synonym should only include the spelled-out terms i.e. "Corynebacterium diphtheriae cross-reacting material 197 protein".
  - Abbreviated organism part names are allowed in a preferred term (and other synonyms). The abbreviations do not need to be accompanied by the fully expanded term, which is an exception to the general naming guidelines on abbreviations and acronyms in the SNOMED CT Editorial Guide. e.g the following PT includes CRM which is the abbreviated form for cross-reacting material.
  - Streptococcus pneumoniae Danish serotype 1 capsular polysaccharide antigen conjugated to Corynebacterium diphtheriae CRM197 protein
• Inclusion of Vaccine manufacturing techniques and/or residuals in antigen names is generally out of scope for the international release. However, exceptions can be made for applicable use-cases.
  - "Antigen of X grown in nervous tissue" or "Antigen of X grown in cellular line" is acceptable for inactivated Rabies antigens grown in brain tissue or cell lines and is included in the international release, considering the difference in the adverse reactions that they cause. If and when created:
    - "Antigen of X grown in cellular line" will be a grouper and other subgroups will be added as needed, e.g. "grown in Human diploid cell".
    - "Antigen of X grown in nervous tissue" should indicate the specific type of nervous tissue e.g. brain.
  - "Split virion" and "surface subunit" is acceptable when the proposed granular antigen is used in a real Clinical Drug, e.g. Influenza antigens.
    - For influenza strains, the candidate vaccine virus (CVV) is not included in the antigen name in the international release. The justification is that each pharmaceutical company may use a different CVV to manufacture their product, or they may omit that level of detail in their product information. However, each of the recommended "parent strain + CVV" are antigenically like the parent virus.
• "Antigen of X adsorbed" is in scope for the international release and is modeled in the Substance hierarchy. It is not required to specify details regarding what is adsorbed with as there is no known use-case at this point.
• With the exception of conjugated proteins/toxoids that are in scope for the international release, inclusion of other adjuvants as well as delivery mechanisms (e.g. Aluminum Salts, Oil-in-Water Emulsions, Virosomes) is currently on hold and will be considered if and when associated requests and uses cases are presented.
• "Purified" will not be included in antigen names as the clinical value of stating "Purified antigen" is not clear. In addition, in some cases, it is implied that an antigen is purified based on the preparation technique. This inconsistency in naming (stated vs. implied) can lead to misclassification. If needed, it can be accomplished by using groupers or other modeling considerations in future. But for now, adding a maintenance burden with very limited benefit does not seem valuable.
• Inclusion of non-antigenic vaccine ingredients such as preservatives and stabilizers is out of scope for the international release.
• For "Antigen of whole inactivated X", the inactivation technique (e.g. heat inactivated, formalin inactivated) is out of scope for antigens in the International Release.

The classification of the antigen variants in the Substance hierarchy:
- Antigen of X
  - Antigen of Live attenuated X
  - Antigen of inactivated whole X
  - Antigen of acellular X
  - Antigen of X [protein/recombinant protein]
  - Antigen of X toxoid
  - Antigen of X polysaccharide/oligosaccharide
  - Antigen of X polysaccharide/oligosaccharide unconjugated
  - Antigen of X polysaccharide/oligosaccharide conjugated
    - Antigen of X polysaccharide/oligosaccharide antigen conjugated to Y

Exemplar
The following illustrates the **stated** and **inferred** view:

Modeling - Antibody to X organism

| Parent concept | Most distal appropriate descendant of [116642004 |Antimicrobial antibody (substance)] |
|----------------|--------------------------------------------------------------------------------|
|                | Exceptions: none
| Semantic tag   | (substance) |
| Definition status | 900000000000074008 |Necessary but not sufficient concept definition status (core metadata concept)|
|                | Exceptions: none

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Naming Guidelines - Antibody to X organism

### FSN

<table>
<thead>
<tr>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Antibody to X organism (substance)</td>
</tr>
<tr>
<td>Example:</td>
</tr>
<tr>
<td>Antibody to Bebaru virus (substance)</td>
</tr>
</tbody>
</table>

### Preferred Term

<table>
<thead>
<tr>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td>X organism antibody</td>
</tr>
<tr>
<td>Example:</td>
</tr>
<tr>
<td>Bebaru virus antibody</td>
</tr>
</tbody>
</table>

### Synonyms

<table>
<thead>
<tr>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td>A synonym that matches FSN</td>
</tr>
<tr>
<td>Example:</td>
</tr>
<tr>
<td>Antibody to Bebaru virus</td>
</tr>
</tbody>
</table>

| Pattern:                                                        |
| X organism Ab                                                   |
| Example:                                                       |
| Bebaru virus Ab                                                  |

| Additional synonyms (e.g. when a legitimate synonyms exist for Organism name) are applicable. |

### Exemplar

The following illustrates the **stated** and **inferred** view:

![Diagram]

**Modeling - Immunoglobulin G, M, A, E, D antibody to X organism**

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of</th>
</tr>
</thead>
<tbody>
<tr>
<td>70095009</td>
<td>Immunoglobulin isotype (substance) and Antibody to X organism (substance)</td>
</tr>
<tr>
<td>Exceptions: none</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(substance)</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Definition status</th>
<th>Necessary but not sufficient concept definition status (core metadata concept)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exceptions: none</td>
<td></td>
</tr>
</tbody>
</table>
Naming Guidelines - Immunoglobulin G, M, A, E, D antibody to X organism

<table>
<thead>
<tr>
<th>FSN</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Immunoglobulin G, M, A, E, D antibody to X organism (substance)</td>
</tr>
<tr>
<td>Example:</td>
<td>• Immunoglobulin M antibody to Clostridium difficile (substance)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• X organism IgG, M, A, E, D</td>
</tr>
<tr>
<td>Example</td>
<td>• Clostridium difficile IgM</td>
</tr>
</tbody>
</table>

| Synonyms       | Pattern: A synonym that matches FSN                                   |
|                | Example                                                                 |
|                | • Immunoglobulin M antibody to Clostridium difficile                   |
| Pattern:       | Anti-X organism IgM                                                   |
| Example        | • Anti-Clostridium difficile IgM                                       |

Additional synonyms (e.g. when a legitimate synonyms exist for Organism name) are applicable.

Exemplar
The following illustrates the **stated** and **inferred** view:

```
716624008
Immunoglobulin M antibody to Clostridium difficile (substance)
```

```
74889000
Immunoglobulin M (substance)
```

```
120950000
Clostridium difficile antibody (substance)
```
Antivenin and descendants
Overview

Modeling

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of 115668003</th>
<th>Biological substance (substance)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Exceptions: none</td>
<td></td>
</tr>
<tr>
<td>Semantic tag</td>
<td>(substance)</td>
<td></td>
</tr>
<tr>
<td>Definition status</td>
<td>900000000000074008</td>
<td>Necessary but not sufficient concept definition status (core metadata concept)</td>
</tr>
<tr>
<td></td>
<td>• Exceptions: none</td>
<td></td>
</tr>
<tr>
<td>Attribute</td>
<td>[Has disposition] = 763291003</td>
<td>Antivenin (disposition)</td>
</tr>
</tbody>
</table>

Naming Guidelines

- Use antivenom, not antivenin, for FSNs and Preferred Terms. Synonyms containing antivenin will not be created routinely, but may be created upon request. Existing concepts that are not consistent with this naming convention will be cleaned up as a batch at a later date.
- FSNs should be based on the scientific name, if there is a one-to-one correspondence. Naming conventions for polyvalent antivenoms (effective against multiple organisms) will not comply with this naming convention, and will be evaluated on a case-by-case basis. Preferred Terms should be based on the common name.
- Synonyms based on the scientific name should be created in most cases.

Useful reference: Current version of *WHO Guidelines for the Production, Control and Regulation of Snake Antivenom Immunoglobulins*

<table>
<thead>
<tr>
<th>FSN</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• X antivenom (substance)</td>
</tr>
<tr>
<td>Example:</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Chironex fleckeri antivenom (substance)</td>
</tr>
</tbody>
</table>
Conjugate acids, bases and salts

Overview
Acids substance concepts should be modeled with a structural parent concept.

Salts should be modeled with a conjugate base as parent (if one exists, otherwise they are modeled with a structural parent) and an Is modification of (attribute) relationship to the conjugate acid.

Conjugate base concepts (e.g. valproate, pamidronate, etidronate) should only be created where a specific use case is identified for example when required to support the definition of other concepts in the terminology.

Conjugate bases should be created as separate concepts - not added as synonyms to the corresponding acid concept.

New instances of substance concepts containing the word "salt" will not be added. The existing concepts will be reviewed and if possible will be replaced by equivalent compounds.

Modeling

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of 312413002 [Substance categorized structurally (substance)]</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Exceptions: none</td>
</tr>
<tr>
<td>Semantic tag</td>
<td>(substance)</td>
</tr>
<tr>
<td>Definition status</td>
<td>90000000000000074008 [Necessary but not sufficient concept definition status (core metadata concept)]</td>
</tr>
<tr>
<td></td>
<td>• Exceptions: none</td>
</tr>
</tbody>
</table>
### Attribute: Is Modification

<table>
<thead>
<tr>
<th>SNOMED CT Concept</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>738774007</td>
<td>Is Modification of</td>
</tr>
</tbody>
</table>

- **Range**: `<105590001 | Substance (substance)`
- **Cardinality**: 0..1
- **Exceptions**: none identified

Used to identify the conjugate acid.

### Attribute: Has disposition

<table>
<thead>
<tr>
<th>SNOMED CT Concept</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>`&lt;726711005</td>
<td>Disposition (disposition)`</td>
</tr>
</tbody>
</table>

**NOTE**: While the allowed range is broader, substance concepts should only use descendants of the concept `<726711005 | Disposition (disposition)` as the attribute value.

- **Cardinality**: 0..1
- **Exceptions**: none identified

### Naming Guidelines

Where INN names exist these should be used for the FSN and PT

#### FSN

**Pattern:**
- X sulfate (substance)
- X pamidronate (substance)

**Example**:
- Copper sulfate (substance)
- Disodium pamidronate (substance)
- Etidronate (substance)

**Exception**:
- Valproate sodium (substance)

#### Preferred Term

**Pattern:**
- X sulfate
- X pamidronate

**Example**:
- Copper sulfate
- Pamidronate monosodium
- Etidronate

**Exception**:
- Valproate sodium

#### Synonyms

**Pattern**: Pamidronate X

**Example**:
- Pamidronate monosodium
- Valproate sodium

### Exemplar

The following illustrates the **stated** and **inferred** views.
The following illustrates the stated and inferred views.
Deoxyribonucleic acid and ribonucleic acid
Modeling - Ribonucleic acid of X organism

| Parent concept | Most distal appropriate descendant of 118248003 |Microbial ribonucleic acid (substance)|
|----------------|---------------------------------------------------|
|                | • Exceptions: none                                |
| Semantic tag   | (substance)                                       |
| Definition status | 900000000000074008 |Necessary but not sufficient concept definition status (core metadata concept)|
|                | • Exceptions: none                                |
| Attribute:     | 738774007 |Is Modification|
| Is Modification | Range: <105590001 |Substance (substance)|
|                | • Cardinality: 0..*                                |
|                | • Exceptions: none identified                      |
|                | • One relationship to be created to represent each modified component of the substance |
| Attribute:     | Range: <726711005 |Disposition (disposition)|
| Has disposition| • NOTE: While the allowed range is broader, substance concepts should only use descendants of the concept 726711005 |Disposition (disposition)| as the attribute value. |
|                | • Cardinality: 0..*                                |
|                | • Exceptions: none identified                      |

Naming Guidelines - Ribonucleic acid of X organism

<table>
<thead>
<tr>
<th>FSN</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Ribonucleic acid of X organism (substance)</td>
</tr>
<tr>
<td>Example:</td>
<td>• Ribonucleic acid of Norovirus genogroup I (substance)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Preferred Term (US/GB), with Initial letter case sensitive:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pattern:</td>
<td>• X organism RNA</td>
</tr>
<tr>
<td>Example:</td>
<td>• Norovirus genogroup I</td>
</tr>
</tbody>
</table>
Synonyms

Pattern: A synonym that matches FSN

Example:

- Ribonucleic acid of Norovirus genogroup I
- Additional synonyms (when a legitimate synonyms exist for Organism name) are applicable.

Exemplar
The following illustrates the stated and inferred view:

Modeling - Ribosomal ribonucleic acid of X organism

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of 118251005</th>
<th>Microbial ribosomal ribonucleic acid (substance)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Exceptions: none</td>
<td></td>
</tr>
</tbody>
</table>

| Semantic tag   | (substance)                                       |                                                 |
| Definition status | 9000000000000074008 | Necessary but not sufficient concept definition status (core metadata concept) |
|                | Exceptions: none                                 |                                                 |

| Attribute: Is Modification | 738774007 | Is Modification |
|                           | Range: <105590001 [Substance (substance)] |
|                           | Cardinality: 0..* |
|                           | Exceptions: none identified |
|                           | One relationship to be created to represent each modified component of the substance |

| Attribute: Has disposition | Range: < 726711005 | Disposition (disposition) |
|                           | NOTE: While the allowed range is broader, substance concepts should only use descendants of the concept 726711005 | Disposition (disposition) as the attribute value. |
|                           | Cardinality: 0..* |
|                           | Exceptions: none identified |

Naming Guidelines - Ribosomal ribonucleic acid of X organism

<table>
<thead>
<tr>
<th>FSN</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Ribosomal ribonucleic acid of X organism (substance)</td>
</tr>
<tr>
<td>Example:</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Ribosomal ribonucleic acid of Candida (substance)</td>
</tr>
</tbody>
</table>
### Preferred Term

Preferred Term, with initial letter case sensitive.

**Pattern:**
- X organism rRNA

**Example**
- Candida rRNA

### Synonyms

**Pattern:**
- A synonym that matches FSN

**Example:**
- Ribosomal ribonucleic acid of Candida
- Additional synonyms (when a legitimate synonyms exist for Organism name) are applicable.

### Exemplar

The following illustrates the **stated** and **inferred** view:

<table>
<thead>
<tr>
<th>Concept</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>707634006</td>
<td>Ribosomal ribonucleic acid of Candida (substance)</td>
</tr>
<tr>
<td>118251005</td>
<td>Ribosomal ribonucleic acid of microorganism (substance)</td>
</tr>
</tbody>
</table>

### Modeling - Messenger ribonucleic acid of X organism

#### Parent concept
Most distal appropriate descendant of 69908008 |Messenger ribonucleic acid (substance)|

- Exceptions: none

#### Semantic tag
(substance)

#### Definition status
900000000000074008 |Necessary but not sufficient concept definition status [core metadata concept]|

- Exceptions: none

#### Attribute: Is Modification

**Range:** <105590001 |Substance (substance)|

**Cardinality:** 0..*

- Exceptions: none identified
- One relationship to be created to represent each modified component of the substance

#### Attribute: Has disposition

**Range:** <726711005|Disposition (disposition)|

- NOTE: While the allowed range is broader, substance concepts should only use descendants of the concept 726711005 [Disposition (disposition)] as the attribute value.

**Cardinality:** 0..*

- Exceptions: none identified

### Naming Guidelines - Messenger ribonucleic acid of X organism
<table>
<thead>
<tr>
<th>FSN</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Messenger ribonucleic acid of X organism (substance)</td>
<td></td>
</tr>
<tr>
<td>Example:</td>
<td></td>
</tr>
<tr>
<td>• Messenger ribonucleic acid of Severe acute respiratory syndrome coronavirus 2 (substance)</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Preferred Term, with Initial letter case sensitive.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pattern:</td>
<td></td>
</tr>
<tr>
<td>• X organism mRNA</td>
<td></td>
</tr>
<tr>
<td>Example</td>
<td></td>
</tr>
<tr>
<td>• SARS-CoV-2 mRNA</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Synonyms</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td>• A synonym that matches FSN</td>
<td></td>
</tr>
<tr>
<td>Example:</td>
<td></td>
</tr>
<tr>
<td>• Messenger ribonucleic acid of Severe acute respiratory syndrome coronavirus 2</td>
<td></td>
</tr>
<tr>
<td>• Additional synonyms (when a legitimate synonyms exist for Organism name) are applicable.</td>
<td></td>
</tr>
</tbody>
</table>

**Exemplar**

![Exemplar diagram](image)

**Modeling - Messenger ribonucleic acid of X organism encoding for specific protein**

| Parent concept | Most distal appropriate descendant of 69908008 |Messenger ribonucleic acid (substance)|
|----------------|-----------------------------------------------|
|                | Exceptions: none                              |

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(substance)</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Definition status</th>
<th>900000000000074008</th>
<th>Necessary but not sufficient concept definition status (core metadata concept)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Exceptions: none</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attribute: Is Modification</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Range &lt;105590001</td>
</tr>
<tr>
<td>• Cardinality: 0..*</td>
</tr>
<tr>
<td>• One relationship to be created to represent each modified component of the substance</td>
</tr>
<tr>
<td>• Exceptions: none identified</td>
</tr>
</tbody>
</table>
Attribute: Has disposition

- Range: <726711005|Disposition (disposition)
- NOTE: While the allowed range is broader, substance concepts should only use descendants of the concept 726711005 |Disposition (disposition) as the attribute value.
- Cardinality: 0..*
- Exceptions: none identified

Naming Guidelines - Messenger ribonucleic acid of X organism encoding for specific protein

<table>
<thead>
<tr>
<th>FSN</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Messenger ribonucleic acid of X organism encoding for protein Y (substance)</td>
</tr>
<tr>
<td>Example</td>
<td>Messenger ribonucleic acid of Severe acute respiratory syndrome coronavirus 2 encoding spike protein (substance)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>X organism mRNA encoding for protein Y</td>
</tr>
<tr>
<td>Example</td>
<td>SARS-CoV-2 mRNA encoding spike protein</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Synonyms</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>A synonym that matches FSN</td>
</tr>
<tr>
<td>Example</td>
<td>Messenger ribonucleic acid of Severe acute respiratory syndrome coronavirus 2 encoding spike protein (substance)</td>
</tr>
<tr>
<td></td>
<td>Additional synonyms (when a legitimate synonyms exist for Organism name) are applicable.</td>
</tr>
</tbody>
</table>

Modeling - Deoxyribonucleic acid of X organism

| Parent concept | Most distal appropriate descendant of 118249006 |Microbial deoxyribonucleic acid (substance)| |
|----------------|-----------------------------------------------|-------------------------------------------|
|                | Exceptions: none                              |

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(substance)</th>
</tr>
</thead>
</table>

| Definition status | 90000000000000074008 |Necessary but not sufficient concept definition status (core metadata concept)| |
|-------------------|---------------------|-----------------------------------------------|
|                    | Exceptions: none    |                                              |
Attribute: Is Modification

738774007 | Is Modification

- Range: <105590001 | Substance (substance)|
- Cardinality: 0..*
- Exceptions: none identified
- One relationship to be created to represent each modified component of the substance

Attribute: Has disposition

- Range: <726711005 | Disposition (disposition)|
- NOTE: While the allowed range is broader, substance concepts should only use descendants of the concept 726711005 | Disposition (disposition)| as the attribute value.
- Cardinality: 0..*
- Exceptions: none identified

Naming Guidelines - Deoxyribonucleic acid of X organism

FSN

Pattern:
- Deoxyribonucleic acid of X organism (substance)
  Example:
  - Deoxyribonucleic acid of Aspergillus terreus (substance)

Preferred Term

Preferred Term (US/GB), with initial letter case sensitive:

Pattern:
- X organism DNA
  Example
  - Aspergillus terreus DNA

Synonyms

Pattern: A synonym that matches FSN

Example:
- Ribonucleic acid of Norovirus genogroup I

- Additional synonyms (when a legitimate synonyms exist for Organism name) are applicable.

Exemplar

The following illustrates the stated and inferred view:

Edible substance and descendants

Overview
Ultimately the intent is to retire this Edible substance subhierarchy since it represents a role. Substance concepts should, where possible, be modeled with either a structural parent or one that denotes source or origin.

**Modeling (stated view)**

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of 762766007</th>
<th>Edible substance (substance)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exceptions: none</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Additional parent concepts may be modeled to denote origin or structural characteristics.</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

**Semantic tag**

(substance)

**Definition status**

900000000000074008 | Necessary but not sufficient concept definition status (core metadata concept) |

Exceptions: none

**Naming Guidelines**

FSN, PT or Synonym should not contain "- dietary" context in descriptions.

FSN, PT or Synonym should not contain brand or trade names.

Culinary name should be used for the FSN and PT when referring to meat products and food. E.g. Beef dripping (substance), Pork sausage (substance).

If there are additional adjectives such as baked, boiled, fried, lean, minced, low-fat, roast, or stewed, these adjectives should proceed the name of source organism, e.g. Fried beef steak (substance), Tinned fish (substance).

**FSN**

Pattern:

- X (substance)
- Example:
  - Betel nut (substance)
  - Wheat dextrin (substance)
  - Tinned fish (substance)

**Preferred Term**

Pattern:

- X
- Example:
  - Betel nut
  - Wheat dextrin
  - Tinned fish

**Synonyms**

Pattern: scientific name if appropriate

Example:

- Areca catechu

**Exemplar**

The following illustrates the **stated** and **inferred** view
Exemplar
The following illustrates the **stated** and **inferred** view

<table>
<thead>
<tr>
<th>Substance concept</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>391730008 Almond oil</td>
<td></td>
</tr>
<tr>
<td>28688002 Fats and oils</td>
<td></td>
</tr>
<tr>
<td>21847005 Oil</td>
<td></td>
</tr>
<tr>
<td>73509005 Salmon</td>
<td></td>
</tr>
<tr>
<td>227110001 Fatty fish</td>
<td></td>
</tr>
</tbody>
</table>

Fractionated and unfractionated substances
Overview
Substance concepts that contain terms such as “fractionated” or “unfractionated” are not created in the Substance hierarchy. "Unfractinated" should not be included when referring to the "entire substance" or "whole substance"; it is implied. "Fractionated" usually refers to a separation technique or process.

Hydrates and anhydrous substances
Modeling (stated view)

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of 105590001 [Substance (substance)]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Semantic tag</td>
<td>(substance)</td>
</tr>
<tr>
<td>Definition status</td>
<td>900000000000074008 [Necessary but not sufficient concept definition status (core metadata concept)]</td>
</tr>
<tr>
<td>Attribute</td>
<td>738774007 [Is modification of (attribute)] relationship to the unspecified substance</td>
</tr>
<tr>
<td></td>
<td>726542003 [Has disposition (attribute)] as appropriate</td>
</tr>
</tbody>
</table>

Modeling (inferred view)
SNOMED CT Editorial Guide
(2021-10-02)

Parent concept
- Exceptions: none

Semantic tag (substance)

Definition status
900000000000074008 |Necessary but not sufficient concept definition status (core metadata concept)|
- Exceptions: none

Attribute
738774007 |Is modification of (attribute)| relationship to the unspecified substance
726542003 |Has disposition (attribute)| as appropriate

Hydrates
For example,
- Ferrous oxalate dihydrate (substance)
- Caffeine hydrate (substance)
- Zinc sulfate heptahydrate (substance)

Hydrates are modeled with an Is modification of (attribute) relationship to the unspecified substance.

Naming Guidelines

FSN
Pattern:
- X hydrate (substance)
- X dihydrate (substance)
- X monohydrate (substance)

Example:
- Copper sulfate pentahydrate (substance)

Preferred Term
Pattern:
- X hydrate
- X dihydrate
- X monohydrate

Example:
- Copper sulfate pentahydrate

Synonyms
Pattern:
- none

Exemplar
The following illustrates the inferred view.
- Caffeine hydrate (substance) IS MODIFICATION of Caffeine (substance)
Anhydrous compounds

Anhydrous compounds have a IS MODIFICATION OF relationship to the unspecified substance

Examples.

Sodium carbonate anhydrous (substance)
Theophylline anhydrous (substance)

Naming Guidelines

FSN Pattern:
- X anhydrous (substance)
- Example:
  - Theophylline anhydrous (substance)

Preferred Term Pattern:
- X anhydrous
  - Example:
    - Theophylline anhydrous

Synonyms Pattern: Anhydrous X
- Example: none

Exemplar

The following illustrates the inferred view.

Theophylline anhydrous (substance) IS MODIFICATION OF Theophylline (substance)
Ions and electrolyte substances

Overview

Previously, SNOMED CT represented ions and electrolytes as separate concepts. Although the two terms are subtly different, there is considerable overlap in their definitions, and they seem to be used interchangeably in medicine. The Substances Redesign Project Group has considered this issue and determined the following:

- Ions should be modeled as child concepts of the substance.
- A second parent (in addition to X (substance)), that indicates the concept is an ionized substance should be modeled.
  - Use a descendant of 86355000 |Electrolyte (substance)| as the additional parent. (86355000 | Electrolyte (substance)| and its children are used in LOINC term expression associations.)
- Non-Ionized/Non-Ionised substances should not be created.

Modeling (stated view)

| Parent concepts | Most distal appropriate descendant of 86355000 |Electrolyte (substance)| |
|-----------------|-------------------------------------------------|-----------------------|
|                 | • Exceptions: none                              |                       |
|                 | Most distal appropriate descendant of 312413002 |Substance categorized structurally (substance)| |
|                 | • Exceptions: none                              |                       |

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(substance)</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Definition status</th>
<th>900000000000000074008</th>
<th>Necessary but not sufficient concept definition status (core metadata concept)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Exceptions: none</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attribute</th>
<th>none</th>
</tr>
</thead>
</table>

Naming Guidelines
### Exemplar

The following illustrates the **Stated** and **Inferred** view

![Exemplar Diagram](image)

The following illustrates the hierarchy view

![Hierarchy Diagram](image)

### Isomers

**Overview**

Within the substances hierarchy in SNOMED CT there are a number of substances where that is either a mix of isomers of a single chemical, a racemic mix or a single isomer.
Each of these substances will all need to be represented in SNOMED CT as separate concepts to support the representation of active ingredients in SNOMED CT and should be represented as sibling concepts without any SNOMED CT relationship created between the two substances.

Exemplar
The following illustrates the hierarchy view for Omeprazole and Esomeprazole:

Metabolites
Overview
Concepts representing Metabolites of X will be considered for inclusion based upon project requirements.

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>767279008 [Metabolite (substance)]</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Exceptions: none</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(substance)</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Definition status</th>
<th>9000000000000074008 [Necessary but not sufficient concept definition status (core metadata concept)]</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Exceptions: none</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attribute</th>
<th>None</th>
</tr>
</thead>
</table>

Naming Guidelines
FSN | Pattern:  
- X metabolite (substance)  
  Example  
  • Cocaine metabolite (substance)

Preferred Term | Pattern:  
- X metabolite  
  Example:  
  • Cocaine metabolite

Synonyms | Pattern:  
- none identified  
  Example:  
  • Cocaine metabolite

Exemplar
The following illustrates the **stated** and **inferred** view:

![Diagram](image)

**Radioactive substances**  
**Representation of radioactive isotopes**  
**Modeling (stated view)**

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of 89457008</th>
<th>Radioactive isotope (substance)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Exceptions: none</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Most distal appropriate descendant of 33638001</td>
<td>Isotope (substance)</td>
</tr>
<tr>
<td></td>
<td>Exceptions: none</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(substance)</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Definition status</th>
<th>9000000000000074008</th>
<th>Necessary but not sufficient concept definition status (core metadata concept)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Exceptions: none</td>
<td></td>
</tr>
</tbody>
</table>
### Attribute: Is Modification

<table>
<thead>
<tr>
<th>ID</th>
<th>Description</th>
<th>Value</th>
<th>Range</th>
<th>Cardinality</th>
<th>Exceptions</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>738774007</td>
<td>Is modification of</td>
<td></td>
<td>&lt;105590001 Substance</td>
<td>0..*</td>
<td>none</td>
<td>One relationship to be created to represent each modified component of actual substance</td>
</tr>
</tbody>
</table>

### Attribute: Has disposition

<table>
<thead>
<tr>
<th>ID</th>
<th>Description</th>
<th>Value</th>
<th>Range</th>
<th>Cardinality</th>
<th>Exceptions</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>726711005</td>
<td>Disposition (disposition)</td>
<td></td>
<td>&lt;726711005 Disposition</td>
<td>0..1</td>
<td>none</td>
<td>While the allowed range is broader, substance concepts should only use descendants of the concept 726711005 Disposition (disposition) as the attribute value.</td>
</tr>
</tbody>
</table>

#### Naming Guidelines

Superscripts should not be used in either Fully Specified Name, Preferred Term, or Synonyms.

**FSN**

<table>
<thead>
<tr>
<th>Pattern</th>
<th>Example</th>
</tr>
</thead>
<tbody>
<tr>
<td>[name of isotope]-[atomic number] (substance)</td>
<td>Gallium-67 (substance)</td>
</tr>
</tbody>
</table>

**Preferred Term**

<table>
<thead>
<tr>
<th>Pattern</th>
<th>Example</th>
</tr>
</thead>
<tbody>
<tr>
<td>[name of isotope]-[atomic number]</td>
<td>Gallium-67</td>
</tr>
</tbody>
</table>

**Synonyms**

<table>
<thead>
<tr>
<th>Pattern</th>
<th>Example</th>
</tr>
</thead>
<tbody>
<tr>
<td>[atomic number]-[chemical symbol]</td>
<td>67-Ga</td>
</tr>
</tbody>
</table>

No synonyms should exist using the superscript notation.

---

**Exemplar**

The following illustrates the hierarchy view:
The following illustrates the **stated** and **inferred** view:

Representation of radioisotope with salt

| Parent concept | Most distal appropriate descendant of 89457008 |Radioactive isotope (substance)|
|----------------|-----------------------------------------------|
| Exceptions:    | none                                          |

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(substance)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Definition status</td>
<td>900000000000074008</td>
</tr>
<tr>
<td>Exceptions:</td>
<td>none</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attribute: Is modification</th>
<th>738774007</th>
<th>Is modification of</th>
</tr>
</thead>
<tbody>
<tr>
<td>Range</td>
<td>&lt;105590001</td>
<td>Substance (substance)</td>
</tr>
<tr>
<td>Cardinality:</td>
<td>0..*</td>
<td></td>
</tr>
<tr>
<td>Exceptions: none</td>
<td>identified</td>
<td></td>
</tr>
<tr>
<td>One relationship</td>
<td>to be created to represent each modified component of the substance</td>
<td></td>
</tr>
</tbody>
</table>

| Attribute: Has disposition | Range: <726711005 |Disposition (disposition)| |
|---------------------------|-------------------|-------------------------|
|  NOTE: While the allowed range is broader, substance concepts should only use descendants of the concept 726711005 as the attribute value. |
|  Cardinality:            | 0..1             |
| Exceptions: none         | identified        |

**Naming Guidelines**

Superscripts should not be used in either Fully Specified Name, Preferred Term, or Synonyms.
Parentheses should be used to delineate the atomic number and chemical symbol from the rest of the terming, this aligns with INN.

<table>
<thead>
<tr>
<th>Terming</th>
<th>Pattern</th>
<th>Example</th>
</tr>
</thead>
</table>
| FSN     | [name of isotope] ([atomic number]-[chemical symbol]) [salt] (substance)  

  - Gallium (67-Ga) citrate (substance) |
| Preferred Term | [name of isotope] ([atomic number]-[chemical symbol]) [salt]  

  - Gallium (67-Ga) citrate |
| Synonyms | none identified |

Exemplar - radioisotope with salt  
The following illustrates the **stated** and **inferred** view:

**Representation of combined radioisotope substances**  
**Modeling (stated view)**

| Parent concept | Most distal appropriate descendant of 89457008 |Radioactive isotope (substance)| and a second parent identifying the labelled component  

  - Exceptions: none |
| Semantic tag | (substance) |
| Definition status | 900000000000074008|Necessary but not sufficient concept definition status (core metadata concept)|  

  - Exceptions: none |
### Attribute

<table>
<thead>
<tr>
<th>Attribute</th>
<th>738774007</th>
<th>Is modification of</th>
</tr>
</thead>
<tbody>
<tr>
<td>Is modification</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

- **Range**: `<105590001 | Substance (substance)`
- **Cardinality**: `0..*
- **Exceptions**: None identified
- **One relationship to be created to represent each modified component of the substance**

| Attribute          | Range: `<726711005 | Disposition (disposition)` |
|--------------------|---------------------|
| Has disposition    |                     |

- **NOTE**: While the allowed range is broader, substance concepts should only use descendants of the concept `<726711005 | Disposition (disposition)` as the attribute value.
- **Cardinality**: `0..1
- **Exceptions**: None identified

### Naming Guidelines

Superscripts should not be used in either Fully Specified Name, Preferred Term, or Synonyms.

The Fully Specified Name should explicitly state that a component was “labeled” by a radioisotope in the FSN. Do not use “with”, “and”, or “tagged”.

Omit the word “labeled” from the PT, keep a synonym to match the FSN.

Exemplar - combined radioisotope substances

The following illustrates both the **stated** and **inferred** view:

- **423249007**: Lactated monomeric antibody (substance)
- **738774007**: Is modification of (attribute)
- **705074002**: Radioisotope labeled monomeric antibody (substance)
- **1139003**: Lactated radioisotope (substance)
- **738774007**: Is modification of (attribute)
- **40837006**: Lactated-124 (substance)
- **738774007**: Is modification of (attribute)
- **49616005**: Monoclonal antibody (substance)

### Saturated or unsaturated substances

#### Overview

**Modeling (stated view)**

| Parent concept          | Most distal appropriate descendant of `115668003 | Biological substance (substance)| |
|-------------------------|-------------------------------------------------|--------------------------------|
|                         | **Exceptions**: None                             |                                |

| Semantic tag            | (substance)                                      |

| Definition status       | 9000000000000074008 | Necessary but not sufficient concept definition status (core metadata concept)| |
|-------------------------|-------------------------------------------------|--------------------------------|
|                         | **Exceptions**: None                             |                                |

| Attribute               | **none**                                         |

### Naming Guidelines
FSN

<table>
<thead>
<tr>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Saturated X (substance)</td>
</tr>
<tr>
<td>• Unsaturated X (substance)</td>
</tr>
<tr>
<td>Example:</td>
</tr>
<tr>
<td>• Unsaturated adipate (substance)</td>
</tr>
</tbody>
</table>

Preferred Term

<table>
<thead>
<tr>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Saturated X</td>
</tr>
<tr>
<td>• Unsaturated X</td>
</tr>
<tr>
<td>Example:</td>
</tr>
<tr>
<td>• Unsaturated adipate</td>
</tr>
</tbody>
</table>

Synonyms

<table>
<thead>
<tr>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td>• X Saturated or X Unsaturated</td>
</tr>
<tr>
<td>Example:</td>
</tr>
<tr>
<td>• Adipate unsaturated</td>
</tr>
</tbody>
</table>

Exemplar
The following illustrates the **stated** and **inferred** view:

![Diagram](image)

**Substances specifying a source or origin**

**Overview**

- A recombinant (A new entity e.g., gene, protein, cell, or individual that results from genetic recombination) or synthetic/semi-synthetic substance (of, relating to, or produced by chemical or biochemical synthesis; especially: produced artificially) is similar in structure (but not identical to the naturally occurring substance). It should be created as a sibling of the naturally occurring substance.
- A substance that is part of (e.g. dander) or is derived from (e.g. Insulin) an organism (human or non-human) is identical and should be created as a child of the naturally occurring substance.

**Modeling**

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of 105590001 [Substance (substance)]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Semantic tag</td>
<td>(substance)</td>
</tr>
<tr>
<td>Definition status</td>
<td>9000000000000074008 [Necessary but not sufficient concept definition status (core metadata concept)]</td>
</tr>
<tr>
<td>- Exceptions: none</td>
<td></td>
</tr>
<tr>
<td>Attribute</td>
<td>[Is modification of] as applicable</td>
</tr>
<tr>
<td></td>
<td>[Has disposition] as appropriate</td>
</tr>
</tbody>
</table>

**Naming Guidelines**
“Recombinant”, “Synthetic”, or “Semi-synthetic” should precede the name of the substance

The source organism name should precede the name of the substance

- An exception to this rule is the naming of DNA, RNA, rRNA, antibody, Immunoglobulin, and antigen of organisms, which should follow the guideline specified for antibodies and antigens
- For some non-human sources, there are different ways of referring to the source organism:
  - The common name of the organism e.g. pig, cow, horse, mouse, sheep
  - The common name of an organism is used when referring to a part of the organism or to non-meat products
    - E.g. Pig epithelium, Cow milk
  - An adjective referring to the organism, e.g. porcine, bovine, equine, murine, ovine
    - The adjective is usually used when referring to a substance extracted from the organism
      - E.g. Bovine growth hormone (substance), Porcine calcitonin (substance)
  - The adjective is commonly used in referring to the organism hosting a virus, but there are exceptions. In general, proper name of organism such as virus name should be preserved as it is (including the case sensitivity):
    - Ribonucleic acid of Bovine leukemia virus (substance)
    - Immunoglobulin G antibody to Eastern equine encephalitis virus (substance)
    - Antigen of Nairobi sheep disease virus (substance)

Additional adjectives (such as labeled, iodinated) should proceed the name of source organism

For example,

- Lente human insulin (substance)
- Iodinated (125-I) human serum albumin (substance)

If a concept includes “recombinant” or “synthetic” as well as a source organism, “recombinant” or “synthetic” should precede the source organism name

For example,

- Recombinant bovine growth hormone (substance)

<table>
<thead>
<tr>
<th>FSN</th>
<th>Pattern:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>[Source organism] X (substance)</td>
</tr>
<tr>
<td></td>
<td>Recombinant X (substance)</td>
</tr>
<tr>
<td></td>
<td>Synthetic X (substance)</td>
</tr>
</tbody>
</table>

Example:

- Human antithrombin III (substance)
- Recombinant thrombin (substance)
- Synthetic steroid (substance)
**Preferred Term**

Pattern:

- [Source organism] X
- Recombinant X
- Synthetic X

Example:

- Human antithrombin III
- Recombinant thrombin
- Synthetic steroid

**Synonyms**

Pattern: none identified

Example:

---

**Exception**

The naming of DNA, RNA, rRNA, antibody, Immunoglobulin, and antigen of organisms should follow the specific guideline for this type of substances.

---

**Exemplar**

Stated view of 411973001 |Bovine thrombin (substance)|

---

**Substances specifying bound**

**Overview**

Creation of substance concepts that contain “bound” is limited to the measurement of biological substances (Observable entities, evaluation procedures, LOINC collaboration project) or to report their level (clinical findings).

Concepts are modeled with an Is modification of (attribute) relationship to the corresponding unspecified substance. Where the substance is "bound" the molecule to which the substance is bound to may be specified in the term.

**Bound substances modeling (stated view)**
Parent concept: Most distal appropriate descendant of 115668003 |Biological substance (substance)|
- Exceptions: none

Semantic tag: (substance)

Definition status: 900000000000074008 |Necessary but not sufficient concept definition status (core metadata concept)|
- Exceptions: none

Attribute: 738774007 |Is modification of (attribute)|
726542003 |Has disposition (attribute)|

Terming Guidelines:

FSN Pattern:
- Bound X (substance)
Example:
- Bound insulin (substance)
- Protein bound iodine (substance)
- Albumin bound thyroxine (substance)

Preferred Term Pattern:
- Bound X (substance)
Example:
- Bound insulin
- Protein bound iodine (substance)
- Albumin bound thyroxine (substance)

Synonyms Pattern:
- X bound
Example:

Exemplar
The following illustrates the stated view:
The following illustrates the **inferred** view:
Substances specifying free or unbound or unconjugated

Overview

Creation of substance concepts that contain terms such as "free" (or "unbound" or "unconjugated") is limited to the measurement of biological substances (Observable entities, evaluation procedures, LOINC collaboration project) or to report their level (clinical findings). These concepts are created as siblings of the base concepts with a "substance X and X derivatives" or "X and X compound" as common supertype.

Modeling (stated view)

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of 105590001 [Substance (substance)]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Semantic tag</td>
<td>(substance)</td>
</tr>
<tr>
<td>Definition status</td>
<td>900000000000074008 [Necessary but not sufficient concept definition status (core metadata concept)]</td>
</tr>
<tr>
<td>Attribute</td>
<td>726542003 [Has disposition (attribute)]</td>
</tr>
</tbody>
</table>

Naming Guidelines

FSN

Pattern: Free X (substance)

Example: Free iodine (substance)

Preferred Term

Pattern: Free X

Example: Free iodine

Synonyms

Pattern: Optional: Unbound iodine

Example:

Exemplar

The following illustrates both the stated and inferred view:

```
706936002
Free iodine (substance)

767266004
Iodine and iodine compound (substance)
```
The following illustrates the hierarchy view:

- Testosterone and testosterone derivative (substance)
- Bioavailable testosterone (substance)

### Substances specifying intact, fragment or subunit

#### Overview
- Fragments to be created as siblings of the intact/whole/entire substance
- "Intact" should not be included when referring to the "entire substance" or "whole substance"; it is implied
- Alpha and beta sub-units should be the children of the entire substance, where "nicked" substances and fragments should be the siblings
- INN should be used for the Preferred Term where they exist

#### Modeling

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of 115668003</th>
<th>Biological substance (substance)</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Exceptions: none</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

| Semantic tag   | (substance)                                    |                                 |                   |
| Definition status | 900000000000000074008 |Necessary but not sufficient concept definition status (core metadata concept)|
|                | • Exceptions: none                             |                                 |                   |

| Attribute      | Has disposition| as appropriate |                   |

#### Naming Guidelines

**FSN**

- **Pattern:**
  - X fragment (substance)
  - X subunit (substance)
  - For example,
    - Corticotrophin big fragment (substance)
    - Thyrotropin beta subunit (substance)
### Preferred Term

<table>
<thead>
<tr>
<th>Pattern:</th>
</tr>
</thead>
</table>
| • X fragment  
  For example,  
  • Corticotrophin big fragment  
  • Thyrotropin beta subunit |

### Synonyms

| Pattern: |
| none identified |

### Substances specifying physical state or physical form

**Overview**

Concepts specifying a particular physical state (e.g. liquid, solid, fumes, vapor, crystal, foam) of a substance should be represented as the sibling for the unspecified substance and also have a parent concept that relates to the structure of the substance.

Concepts specifying a particular physical form (e.g. micronized, macrocrystal, microsphere) should have a parent concept that relates to the structure of the substance and also an Is modification of (attribute) relationship to the unspecified substance concept.

### Modeling - Micronized substance

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>Most distal appropriate descendant of 105590001 [Substance (substance)]</th>
</tr>
</thead>
</table>
| • Exceptions: none  
  An additional parent concept to be allocated to denote physical form as appropriate |

| Semantic tag | (substance) |

<table>
<thead>
<tr>
<th>Definition status</th>
<th>900000000000074008 [Necessary but not sufficient concept definition status (core metadata concept)]</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Exceptions: none</td>
<td></td>
</tr>
</tbody>
</table>

| Attribute | 738774007 [Is modification of (attribute)] relationship to the unspecified substance  
  726542003 [Has disposition (attribute)] as appropriate |

### Naming Guidelines

Fumes should be expressed in the plural (i.e. fumes as opposed to fume)

Fumes and vapor are different and so should not be used as synonyms on the same concept [https://www.commerce.wa.gov.au/worksafe/gases-vapours-smoke-and-fumes](https://www.commerce.wa.gov.au/worksafe/gases-vapours-smoke-and-fumes)

### FSN

<table>
<thead>
<tr>
<th>Pattern:</th>
</tr>
</thead>
</table>
| • X fumes (substance)  
  • X micronized (substance)  
  Example  
  • Bauxite fumes (substance)  
  • Fenofibrate micronized (substance) |
### Preferred Term

**Pattern:**
- X fumes
- X micronized

**Example:**
- Bauxite fumes
- Fenofibrate micronized

### Synonyms

**Pattern:** X form to represent EN-GB language variants where appropriate

**Example**
- Fenofibrate micronised

---

**Exemplar**

Figure: View of both stated and inferred form of 278945005 |Iron fumes (substance)|
Appendices

Editorial Guide Style and Terms

To provide consistency and clarity, there has been an effort to use certain styles and specific terms within the Editorial Guide. Although this has been the intent, it is the content of the Guide that has been the focus. Consequently, authors may find instances where alternative styles or terms are used.

<table>
<thead>
<tr>
<th>Style</th>
<th>Notes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Italics</td>
<td>• To emphasize a word in a sentence or phrase</td>
</tr>
<tr>
<td></td>
<td>• To indicate the name of something</td>
</tr>
<tr>
<td>Style</td>
<td></td>
</tr>
<tr>
<td>-------</td>
<td></td>
</tr>
<tr>
<td><strong>Upper case first letter</strong></td>
<td></td>
</tr>
<tr>
<td>• To emphasize a word in a sentence or phrase not necessarily at the beginning</td>
<td></td>
</tr>
</tbody>
</table>

| Periods. |
| Not used in: |
| • Lists when items contained therein are not sentences |
| • At the end of sentences within tables |

| Examples |
| • Presented as: |
| For example, |
| • text of example |
| • When possible, examples from the SNOMED CT browser are provided. When examples from the browser are not available, i.e do not yet exist, they are obtained from other sources |

| Macros: |
| • Note = yellow |
| • For example |
| • For more information |
| • Tip = green |
| • For example |
| • Concept modeling |
| • URLs |
| • Warning = red |
| • Exceptions |
| • Info = blue |

| General |
| **For more information** |
| Contains general information or additional resource (may be a link) |

| **Modeling** |
| Contains modeling information |

| **Exception or Inactivation or Under Revision** |
| Contains information about content exceptions, inactivation, or under revision |

| **Hello** |
| Informational box |

| Tables |
| • Heading row - light green fill; **bold font** |
| • Sub-heading row - light blue fill |
| • Column heading - light blue fill |
| • Other cells - may have pink fill for emphasis |
| • Footing row - light yellow fill |

| **Heading row** |

| **Footing row** |

| **Note:** Tables generated from the Human Readable Concept Model have unique formatting |

| Text formats |
| • "Quotation marks" |
| • **Bold font** |
| • ALL CAPS |

<p>| • Minimally used |</p>
<table>
<thead>
<tr>
<th><strong>Style</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Page headings</strong></td>
</tr>
<tr>
<td>• Section headings - Heading 2</td>
</tr>
<tr>
<td>• Subsection headings - Heading 3</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th><strong>Terms</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Used in Guide</strong></td>
</tr>
<tr>
<td>Synonym / Other</td>
</tr>
<tr>
<td><strong>Attribute</strong></td>
</tr>
<tr>
<td>Concept Model Attribute; Relation type</td>
</tr>
<tr>
<td><strong>Authoring platform</strong></td>
</tr>
<tr>
<td>SCA tool</td>
</tr>
<tr>
<td><strong>Child</strong></td>
</tr>
<tr>
<td>Subtype, Subtype child</td>
</tr>
<tr>
<td><strong>Classifier</strong></td>
</tr>
<tr>
<td>Description logic (DL) classifier; Logic reasoner</td>
</tr>
<tr>
<td><strong>Descendant</strong></td>
</tr>
<tr>
<td>Child and Subtype child/children</td>
</tr>
<tr>
<td><strong>Domain</strong></td>
</tr>
<tr>
<td>Concept model domain</td>
</tr>
<tr>
<td><strong>Electronic health application</strong></td>
</tr>
<tr>
<td>Software application; Clinical information systems</td>
</tr>
<tr>
<td><strong>Electronic health record or EHR</strong></td>
</tr>
<tr>
<td>Electronic medical record; Electronic record; Electronic patient record</td>
</tr>
<tr>
<td><strong>Extension</strong></td>
</tr>
<tr>
<td>SNOMED CT extension, Member/Affiliate-Added Terminology</td>
</tr>
<tr>
<td><strong>Grouping concept/Grouper</strong></td>
</tr>
<tr>
<td>NA</td>
</tr>
<tr>
<td><strong>Inactivate/Inactivation</strong></td>
</tr>
<tr>
<td>Retire*</td>
</tr>
<tr>
<td><strong>International Release</strong></td>
</tr>
<tr>
<td>Core</td>
</tr>
<tr>
<td><strong>Material entity</strong></td>
</tr>
<tr>
<td>Material agent</td>
</tr>
<tr>
<td><strong>Modeler/Modeling</strong></td>
</tr>
<tr>
<td>Author/Authoring, Editor/Editing</td>
</tr>
<tr>
<td><strong>Parent</strong></td>
</tr>
<tr>
<td>Supertype, Supertype parent</td>
</tr>
<tr>
<td><strong>Precoordinated</strong></td>
</tr>
<tr>
<td>Precoordinated expression</td>
</tr>
<tr>
<td><strong>Postcoordinated</strong></td>
</tr>
<tr>
<td>Postcoordinated expression</td>
</tr>
<tr>
<td><strong>Qualifier</strong></td>
</tr>
<tr>
<td>Qualifying characteristic</td>
</tr>
<tr>
<td><strong>Range</strong></td>
</tr>
<tr>
<td>Concept model range, Allowable value</td>
</tr>
<tr>
<td><strong>Relationship group</strong></td>
</tr>
<tr>
<td>Role group</td>
</tr>
<tr>
<td><strong>Root concept</strong></td>
</tr>
<tr>
<td>Top-level concept</td>
</tr>
<tr>
<td><strong>Semantic tag</strong></td>
</tr>
<tr>
<td>Semantic type, Hierarchy tag, Hierarchy designator</td>
</tr>
<tr>
<td><strong>Situation with Explicit Context</strong></td>
</tr>
<tr>
<td>Context-dependent Category</td>
</tr>
<tr>
<td><strong>Sufficiently defined</strong></td>
</tr>
<tr>
<td>Fully-defined*</td>
</tr>
</tbody>
</table>

*Strikethrough = No longer used
SNOMED CT Editorial Guide - Pharmaceutical Dose Form

Editorial guidelines for the Pharmaceutical dose form hierarchy and for supporting hierarchies required to support creation of sufficiently defined pharmaceutical dose form concepts are documented in the following sections.

Table of Contents

Summary of Changes

<table>
<thead>
<tr>
<th>Date</th>
<th>Comments</th>
<th>Effective date</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Added Appendix A - Cumulative Summary of Changes</td>
<td>2021-Jul</td>
</tr>
</tbody>
</table>

Assumptions, Requirements, and Scope

General Assumptions and Requirements

<table>
<thead>
<tr>
<th>Assumption or Requirement</th>
</tr>
</thead>
<tbody>
<tr>
<td>1  Concept model will conform to description logic principles, including use of the classifier to organize the concepts in the hierarchy.</td>
</tr>
<tr>
<td>2  Concept model will include attributes necessary to define the concepts to ensure consistent and reproducible modeling of concepts, whose use is primarily to describe or group concepts in the Medicinal Product hierarchy.</td>
</tr>
</tbody>
</table>
| 3  Any requirement to align to external standards or registries will be explicitly documented.  
   Concept model will be compatible with the following ISO (International Organization for Standardization) IDMP (Identification of Medicinal Products) standards where appropriate:  
   • ISO 11239 Health informatics — Identification of medicinal products — Data elements and structures for the unique identification and exchange of regulated information on pharmaceutical dose forms, units of presentation, routes of administration and packaging  
   • ISO/TS 20440 Health informatics — Identification of medicinal products — Implementation guide for ISO 11239 data elements and structures for the unique identification and exchange of regulated information on pharmaceutical dose forms, units of presentation, routes of administration and packaging |
| 4  Concepts in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy shall be sufficiently defined using proximal primitive modeling methodology unless explicitly noted as an exception in the editorial guidelines.  
   • Additional information about proximal primitive modeling and intermediate primitive concepts can be found in the main SNOMED CT Editorial Guide. |
| 5  Content in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy in the International Release is not intended to eliminate the need for a national extension. |

Scope

<table>
<thead>
<tr>
<th>Scope</th>
</tr>
</thead>
<tbody>
<tr>
<td>1  Concepts representing combined dose forms are out of scope (e.g. single concepts describing the multiple dose forms found in kit products such as cream and pessary).</td>
</tr>
<tr>
<td>2  Concepts representing proprietary dose forms are out of scope.</td>
</tr>
</tbody>
</table>
Concepts that are not allowed to be used in modeling Medicinal product concepts in the International Release may be added to the Pharmaceutical dose form hierarchy to support national extension modeling (e.g. 420378007 |Prolonged-release film-coated oral tablet (dose form)|).

### Naming and Modeling Guidelines

#### Table of Contents

**Overview**

The 736542009 |Pharmaceutical dose form (dose form)| hierarchy is comprised of the types of concepts as shown in the table below. Detailed editorial guidelines for each distinct concept type, including required attributes and naming guidelines, are found in the sections that follow.

For the purposes of the following editorial guidelines, pharmaceutical dose form refers to the physical manifestation of a medicinal product that contains the active ingredient substance(s) and inactive ingredient substances that are intended for administration for the patient.

<table>
<thead>
<tr>
<th>Concept type</th>
<th>Example</th>
</tr>
</thead>
<tbody>
<tr>
<td>Grouper based on intended site</td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td>• 740596000</td>
</tr>
<tr>
<td></td>
<td>• 385268001</td>
</tr>
<tr>
<td>Grouper concept without basic dose form</td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td>• 385105007</td>
</tr>
<tr>
<td></td>
<td>• 385136004</td>
</tr>
<tr>
<td>Pharmaceutical dose form</td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td>• 385151008</td>
</tr>
<tr>
<td></td>
<td>• 421026006</td>
</tr>
<tr>
<td></td>
<td>• 385053008</td>
</tr>
</tbody>
</table>

### Grouper Based on Intended Site

#### Overview

Pharmaceutical dose form grouper concepts based on intended site of use for the dose form that are deemed to be clinically useful or which provide a helpful organizing grouper and that can be sufficiently defined may be included in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy.

These concepts are used in modeling (medicinal product form) concepts in the International Release; they are not allowed to model (clinical drug) concepts in the International Release.

#### Modeling (stated view)

Pharmaceutical dose form grouper concepts based on intended site shall be modeled using the proximal primitive modeling pattern.

| Parent concept                  | 736542009 |Pharmaceutical dose form (dose form)| |
|---------------------------------|-----------------------------------------------|
|                                 | • Exceptions: none                            |
### Semantic tag
(dose form)

### Definition status
900000000000073002 |Sufficiently defined by necessary conditions definition status (core metadata concept)|
- Exceptions: none

### Attribute:
#### Has dose form intended site
- Range: << 736479009|Dose form intended site (intended site)
- Cardinality: 0..*
  - NOTE: While the allowed range is broader, the grouper concepts based on dose form intended site should have one or more |Has dose form intended site| attributes.
- Exceptions: none

### Naming Guidelines

**FSN**
Use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concept that is selected as the attribute value, excluding the semantic tag. For multiple intended sites, the intended sites must be in alphabetical order and separated by the word "or".
- <Dose form intended site FSN> dose form (dose form)
- <Dose form intended site FSN> or <Dose form intended site FSN> dose form (dose form)

Example:
- Cutaneous dose form (dose form)
- Oral dose form (dose form)
- Parenteral dose form (dose form)
- Ocular or otic dose form (dose form)
- Nasal or ocular or otic dose form (dose form)

**Preferred Term**
Use the following pattern for the PT; align naming and case sensitivity with the PT for the concept that is selected as the attribute value. For multiple intended sites, the intended sites must be in alphabetical order and separated by the word "or".
- <Dose form intended site PT> dose form (dose form)
- <Dose form intended site PT> or <Dose form intended site PT> dose form (dose form)

Example:
- Cutaneous dose form
- Oral dose form
- Parenteral dose form
- Ocular or otic dose form
- Nasal or ocular or otic dose form

**Synonyms**
Synonyms are not allowed unless explicitly identified as an exception in the Editorial Guidelines.
- Exceptions: none identified

**Text definitions**
Optional

### Exemplars
The following illustrates the stated view for grouper concept 740596000 |Cutaneous dose form (dose form)|:
The following illustrates the **inferred** view for grouper concept 740596000 |Cutaneous dose form (dose form)|:

- **740596000** |Cutaneous dose form (dose form) |
- **736542000** |Pharmaceutical dose form (dose form) |
- **736474004** |Has dose form intended site (attribute) |
- **738904002** |Cutaneous (intended site) |

The following illustrates the **stated** view for grouper concept 765166009 |Ocular or otic dose form (dose form)|:

- **765166009** |Ocular or otic dose form (dose form) |
- **736542000** |Pharmaceutical dose form (dose form) |
- **736474004** |Has dose form intended site (attribute) |
- **738952007** |Ocular (intended site) |
- **738983006** |Otic (intended site) |

The following illustrates the **inferred** view for grouper concept 765166009 |Ocular or otic dose form (dose form)|:
Grouper Without Basic Dose Form

Overview
Pharmaceutical dose form grouper concepts that do not include a basic dose form but are deemed to be clinically useful and that can be sufficiently defined will be included in the 736542009 [Pharmaceutical dose form (dose form)] hierarchy. These concepts are particularly useful when there is no clinical need to describe whether the basic dose form is a solution, a suspension or an emulsion (e.g. for products administered to the eyes or nose in the form of drops or spray).

Modeling (stated view)
Grouper concepts concepts that do not include a basic dose form shall be modeled using the proximal primitive modeling pattern.

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>736542009 [Pharmaceutical dose form (dose form)]</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Exceptions: none</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(dose form)</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Definition status</th>
<th>900000000000073002 [Sufficiently defined by necessary conditions definition status (core metadata concept)]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exceptions:</td>
<td>- Grouper concepts representing drug delivery systems will have Definition status = 900000000000074008 [Not sufficiently defined by necessary conditions definition status (core metadata concept)]</td>
</tr>
<tr>
<td>Attribute:</td>
<td></td>
</tr>
<tr>
<td>-----------------------------------------------</td>
<td>-----------------------------------------------------------------------------------</td>
</tr>
<tr>
<td>Has dose form release characteristic</td>
<td>• Range: &lt; 736480007</td>
</tr>
<tr>
<td></td>
<td>• Cardinality: 0..*</td>
</tr>
<tr>
<td></td>
<td>• NOTE: While the allowed range is broader, the grouper concepts without basic dose form should have 1..1</td>
</tr>
<tr>
<td></td>
<td>• Exceptions: none</td>
</tr>
<tr>
<td>Attribute:</td>
<td></td>
</tr>
<tr>
<td>Has dose form intended site</td>
<td>• Range: &lt; 736479009</td>
</tr>
<tr>
<td></td>
<td>• Cardinality: 0..*</td>
</tr>
<tr>
<td></td>
<td>• NOTE: While the allowed range is broader, the grouper concepts without basic dose form should have 1..*</td>
</tr>
<tr>
<td></td>
<td>• Exceptions: none</td>
</tr>
<tr>
<td>Attribute:</td>
<td></td>
</tr>
<tr>
<td>Has dose form administration method</td>
<td>• Range: &lt; 736665006</td>
</tr>
<tr>
<td></td>
<td>• Cardinality: 0..*</td>
</tr>
<tr>
<td></td>
<td>• NOTE: While the allowed range is broader, the grouper concepts without basic dose form should have 1..1</td>
</tr>
<tr>
<td></td>
<td>• Exceptions: none</td>
</tr>
</tbody>
</table>
## Naming Guidelines

| FSN | For concepts with 736472000 [Has dose form administration method (attribute)] = 738996007 | Spray (administration method)], use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concept that is selected as the attribute value. For multiple intended sites, the intended sites must be in alphabetical order and separated by the word “or”.

- <Dose form release characteristic> <Dose form intended site FSN> <Dose form administration method> (dose form)

  Example:
  - Conventional release cutaneous spray (dose form)
  - Conventional release nasal spray (dose form)
  - Conventional release sublingual spray (dose form)

| | For concepts representing drops with 736472000 [Has dose form administration method (attribute)] = 738994005 | Instill (administration method)], use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concept that is selected as the attribute value. For multiple intended sites, the intended sites must be in alphabetical order and separated by the word “or”.

- <Dose form release characteristic> <Dose form intended site FSN> <Dose form administration method> (dose form)

  Example:
  - Conventional release nasal drops (dose form)
  - Prolonged-release eye drops (dose form)

| | For concepts representing drug delivery systems, use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concept that is selected as the attribute value. For multiple intended sites, the intended sites must be in alphabetical order and separated by the word “or”.

- <Dose form release characteristic> <Dose form intended site FSN> drug delivery system

  Example:
  - Prolonged-release intrauterine drug delivery system (dose form)
  - Prolonged-release transdermal drug delivery system (dose form) |
**Preferred Term**

For concepts with 736472000 |Has dose form administration method (attribute)| = 738996007 | Spray (administration method)|, use the following pattern for the PT; align naming and case sensitivity with the PT for the concept that is selected as the attribute value. For multiple intended sites, the intended sites must be in alphabetical order and separated by the word “or”. Exclude <Dose form release characteristic> when = 736849007 |Conventional release (release characteristic)|.

- **<Dose form release characteristic> <Dose form intended site FSN> <Dose form administration method>**

Example:

- Cutaneous spray
- Nasal spray
- Sublingual spray

---------------

For concepts representing drops with 736472000 |Has dose form administration method (attribute)| = 738994005 |Instill (administration method)|, use the following pattern for the PT; align naming and case sensitivity with the PT for the concept that is selected as the attribute value. For multiple intended sites, the intended sites must be in alphabetical order and separated by the word “or”. Exclude <Dose form release characteristic> when = 736849007 |Conventional release (release characteristic)|.

- **<Dose form release characteristic> <Dose form intended site FSN> <Dose form administration method>**

Example:

- Nasal drops
- Prolonged-release eye drops

---------------

For concepts representing drug delivery systems, use the following pattern for the PT; align naming and case sensitivity with the PT for the concept that is selected as the attribute value. For multiple intended sites, the intended sites must be in alphabetical order and separated by the word “or”. Exclude <Dose form release characteristic> when = 736849007 |Conventional release (release characteristic)|.

- **<Dose form release characteristic> <Dose form intended site FSN> drug delivery system**

Example:

- Prolonged-release intrauterine drug delivery system (dose form)
- Prolonged-release transdermal drug delivery system (dose form)

**Synonyms**

A synonym matching the FSN is required; additional synonyms are not allowed unless explicitly identified as an exception in the Editorial Guidelines.

Exceptions:

- Synonyms with eye instead of ocular, ear instead of otic, or nose instead of nasal may be created

**Text definitions**

Optional

**Exemplars**

The following illustrates the **stated** view for concept 385105007 |Conventional release cutaneous spray (dose form)|:
The following illustrates the inferred view for concept 385105007 |Conventional release cutaneous spray (dose form)|:

The following illustrates the stated view for concept 385152001 |Conventional release nasal drops (dose form)|:

The following illustrates the inferred view for concept 385152001 |Conventional release nasal drops (dose form)|:
The following illustrates the **stated** view for concept 421716009 |Prolonged-release transdermal drug delivery system (dose form)|:

The following illustrates the **inferred** view for concept 421716009 |Prolonged-release transdermal drug delivery system (dose form)|:
Pharmaceutical Dose Form

Overview
Pharmaceutical dose form concepts (e.g. conventional release oral tablet, prolonged-release oral capsule) that are deemed to be clinically useful and that can be sufficiently defined will be included in the 736542009 | Pharmaceutical dose form (dose form) hierarchy. Primitive concepts may be included if documented as an exception.

Modeling (stated view)
Pharmaceutical dose form concepts shall be modeled as follows:

<p>| Parent concept | 736542009|Pharmaceutical dose form (dose form) |
|----------------|--------------------------------------------------|
|                | • Exceptions: none |
| Semantic tag   | (dose form) |</p>
<table>
<thead>
<tr>
<th>Definition status</th>
<th>900000000000073002</th>
<th>Sufficiently defined by necessary conditions definition status (core metadata concept)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exceptions:</td>
<td></td>
<td>• The following concepts cannot be sufficiently defined and will be modeled with the parent concept 736542009</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Concepts that refer to &quot;aerosol&quot; (e.g. 420407000</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Concepts that refer to &quot;coated&quot; (e.g. 421338009</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Concepts that refer to &quot;drug delivery system&quot; (e.g. 784576003</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Concepts that refer to &quot;iontophoresis&quot; (e.g. 385113008</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Concepts that refer to &quot;liposomal suspension&quot; (e.g. 420656008</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Concepts that refer to &quot;nebulizer&quot; (e.g. 385198000</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Concepts that refer to &quot;pellet&quot; or &quot;particle&quot; (e.g. 420767002</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Concepts that refer to &quot;syrup&quot; (e.g. 385033009</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• Concepts that refer to &quot;vapor&quot; (e.g. 385215007</td>
</tr>
</tbody>
</table>

| Attribute:        | Has basic dose form | • Range: 736478001 | Basic dose form (basic dose form) - descendants only |
|                  |                     | • Cardinality: 0..1 |
|                  |                     | • NOTE: While the allowed range is broader, concepts representing a sufficiently defined pharmaceutical dose form should have one and only one | Has basic dose form | attributes. |
|                  |                     | • Exceptions: none |
| Attribute: Has dose form intended site | • Range: 736479009 | Dose form intended site (intended site) - descendants only  
| | • Cardinality: 0..*  
| | • NOTE: While the allowed range is broader, concepts representing a sufficiently defined pharmaceutical dose form should have one or more |Has dose form intended site| attributes.  
| | • Exceptions:  
| | 785898006 | Conventional release solution for irrigation (dose form)|  
| | 785910004 | Prolonged-release intralesional implant (dose form)| |
| Attribute: Has dose form release characteristic | • Range: 736480007 | Dose form release characteristic (release characteristic) - descendants only  
| | • Cardinality: 0..1  
| | • NOTE: While the allowed range is broader, concepts representing a sufficiently defined pharmaceutical dose form should have one and only one |Has dose form release characteristic| attributes.  
| | • Exceptions: none |
| Attribute: Has dose form administration method | • Range: 736665006 | Dose form administration method (administration method) - descendants only  
| | • Cardinality: 0..*  
| | • NOTE: While the allowed range is broader, concepts representing a sufficiently defined pharmaceutical dose form should have one and only one |Has dose form administration method| attributes.  
| | • Exceptions: none |
| Attribute: Has dose form transformation | • Range: 736477006 | Dose form transformation (transformation) - descendants only  
| | • Cardinality: 0..*  
| | • NOTE: While the allowed range is broader, concepts representing a sufficiently defined pharmaceutical dose form should have one and only one |Has dose form transformation| attributes.  
| | • Exceptions: none |

Naming Guidelines
### FSN

Use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concepts that are selected as the attribute values, excluding the semantic tag. For multiple intended sites, the sites must be in alphabetical order and separated by the word “or”.

- `<Dose form release characteristic FSN> <Dose form intended site FSN> <Basic dose form> (dose form)`

**Example:**

- Conventional release oral capsule (dose form)
- Conventional release oral suspension (dose form)
- Prolonged-release oral capsule (dose form)
- Conventional release and prolonged-release oral tablet (dose form)
- Conventional release cutaneous cream (dose form)
- Conventional release vaginal ointment (dose form)
- Gastro-resistant oral suspension (dose form)

### Preferred Term

Use the following pattern for the PT; align naming and case sensitivity with the PT for the concepts that are selected as the attribute values, excluding the semantic tag. For multiple intended sites, the sites must be in alphabetical order and separated by the word “or”. Exclude `<Dose form release characteristic>` when = 736849007 [Conventional release (release characteristic)].

- `<Dose form release characteristic FSN> <Dose form intended site FSN> <Basic dose form>`

**Example:**

- Oral capsule
- Oral suspension
- Prolonged-release oral capsule
- Conventional release and prolonged-release oral tablet
- Cutaneous cream
- Vaginal ointment
- Gastro-resistant oral suspension

### Synonyms

A synonym matching the FSN is required; other synonyms are not allowed unless explicitly identified as an exception in the Editorial Guidelines.

**Exceptions:**

- Synonyms with eye instead of ocular, ear instead of otic, or nose instead of nasal may be created

### Text Definitions

Optional

### Exemplar

The following illustrates the **stated** view for 385024007 [Conventional release oral suspension (dose form)]:
The following illustrates the **inferred** view for 385024007 |Conventional release oral suspension (dose form)|:

The following illustrates the **stated** view for 764774009 |Conventional release and prolonged-release oral tablet (dose form)|:
Supporting Hierarchies

Editorial guidelines for the supporting hierarchies required to support creation of sufficiently defined pharmaceutical dose form concepts are documented in the following sections.

Table of Contents
Basic Dose Form Hierarchy

Overview

The basic dose form represents a general type of pharmaceutical formulation (e.g. tablet, capsule, cream, ointment, solution, emulsion) used for medicinal products. To support fully defining concepts in the 736542009 | Pharmaceutical dose form (dose form)| hierarchy, a hierarchy representing basic dose form is required.

Concepts in the 736478001 |Basic dose form (basic dose form)| hierarchy will be used to model concepts in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy; they will not be used to model concepts in the 763158003 |Medicinal product (product)| hierarchy.

The 736478001 |Basic dose form (basic dose form)| hierarchy is a descendant of 362981000 |Qualifier value (qualifier value)|.

Modeling (stated view)

Descendants shall be modeled as follows.

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>736478001</th>
<th>Basic dose form (basic dose form)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Semantic tag</td>
<td>(basic dose form)</td>
<td></td>
</tr>
<tr>
<td>Definition status</td>
<td>900000000000074008</td>
<td>Necessary but not sufficient concept definition status (core metadata concept)</td>
</tr>
<tr>
<td></td>
<td>Exceptions: none</td>
<td></td>
</tr>
<tr>
<td>Attribute:</td>
<td>Has state of matter (attribute)</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Range: 736471007</td>
<td>State of matter (state of matter) - descendants only</td>
</tr>
<tr>
<td></td>
<td>Cardinality: 1..1</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Exceptions: none</td>
<td></td>
</tr>
</tbody>
</table>

Naming Guidelines for Grouper Concept

<table>
<thead>
<tr>
<th>FSN</th>
<th>Use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concept that is selected as the attribute value, excluding the semantic tag.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Basic dose form with <code>&lt;State of matter&gt;</code> state of matter (basic dose form)</td>
</tr>
<tr>
<td>Example:</td>
<td>• Basic dose form with liquid state of matter (basic dose form)</td>
</tr>
<tr>
<td></td>
<td>• Basic dose form with solid state of matter (basic dose form)</td>
</tr>
</tbody>
</table>
Preferred Term

Use the following pattern for the PT; align naming and case sensitivity with the PT for the concept that is selected as the attribute value.

- `<State of matter PT> state of matter`
  
  Example:
  
  - Liquid dose form
  - Solid dose form

Synonyms

A synonym to match the FSN is required.

Additional synonyms are not allowed unless explicitly identified as an exception in the Editorial Guidelines.

- Exceptions: none

Text Definitions

Preferred; not required.

Exemplars for Grouper Concept

The following illustrates the **stated** view for 420699003 |Basic dose form with liquid state of matter (basic dose form)|:

![Diagram](image1)

The following illustrates the **inferred** view for 420699003 |Basic dose form with liquid state of matter (basic dose form)|:

![Diagram](image2)
## Naming Guidelines for Basic Dose Form Concept

<table>
<thead>
<tr>
<th></th>
<th>Use the following pattern for the FSN where X is the basic dose form:</th>
</tr>
</thead>
<tbody>
<tr>
<td>FSN</td>
<td>X (basic dose form)</td>
</tr>
<tr>
<td>Example</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Cream (basic dose form)</td>
</tr>
<tr>
<td></td>
<td>• Gel (basic dose form)</td>
</tr>
<tr>
<td></td>
<td>• Suppository (basic dose form)</td>
</tr>
<tr>
<td></td>
<td>• Tablet (basic dose form)</td>
</tr>
<tr>
<td>Exceptions:</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Plural form to be used for Granules (basic dose form)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th>Use the following pattern for the PT where X is the basic dose form:</th>
</tr>
</thead>
<tbody>
<tr>
<td>Preferred Term</td>
<td></td>
</tr>
<tr>
<td></td>
<td>X</td>
</tr>
<tr>
<td>Example</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Cream</td>
</tr>
<tr>
<td></td>
<td>• Gel</td>
</tr>
<tr>
<td></td>
<td>• Suppository</td>
</tr>
<tr>
<td></td>
<td>• Tablet</td>
</tr>
<tr>
<td>Exceptions:</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Plural form to be used for Granules</td>
</tr>
</tbody>
</table>

### Synonyms

Synonyms are not allowed unless explicitly identified as an exception in the Editorial Guidelines.

- Exceptions: none

### Text Definitions

Text definitions are not required but are encouraged.

---

### Exemplars for Basic Dose Form Concept

The following illustrates the **stated** view for 739006009 [Solution (basic dose form)]:

![Diagram illustrating the stated view for Solution (basic dose form)](attachment:diagram.png)

The following illustrates the **inferred** view for 739006009 [Solution (basic dose form)]:

![Diagram illustrating the inferred view for Solution (basic dose form)](attachment:diagram.png)
Dose Form Administration Method Hierarchy

Overview

The dose form administration method represents a general type of method of administration (e.g. apply, chew, spray, swallow) that a dose form is designed to be administered by (e.g. a chewable tablet is formulated to be administered by chewing). To support fully defining concepts in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy, a hierarchy representing dose form administration method is required.

Concepts in the 736665006 |Dose form administration method (administration method)| hierarchy will be used to model concepts in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy; they will not be used to model concepts in the 763158003 |Medicinal product (product)| hierarchy.

The 736665006 |Dose form administration method (administration method)| hierarchy is a descendant of 362981000 |Qualifier value (qualifier value)|.

Modeling (stated view)

Descendants shall be modeled as follows.

<table>
<thead>
<tr>
<th>Parent concept</th>
<th>736665006</th>
<th>Dose form administration method (administration method)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Exceptions: none</td>
<td></td>
</tr>
</tbody>
</table>

| Semantic tag                                         | (administration method)                             |

| Definition status                                    | 900000000000074008 | Necessary but not sufficient concept definition status (core metadata concept) |
|                                                      | • Exceptions: none                                   |

| Attributes                                           | None                                                |

Naming Guidelines

| FSN | Use the following pattern for the FSN where X is the administration method and is in the form of an imperative:  
|     | X (administration method)                           |
|     | Example:                                            |
|     | • Administer (administration method)                |
|     | • Apply (administration method)                     |
|     | • instill (administration method)                   |
|     | • Spray (administration method)                     |
|     | • Swallow (administration method)                   |

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### Preferred Term

Use the following pattern for the PT where X is the administration method:

\[X\]

Example:
- Administer
- Apply
- Instill
- Spray
- Swallow

### Synonyms

Synonyms are not allowed unless explicitly identified as an exception in the Editorial Guidelines.

Exceptions: none

### Text Definitions

Required.

### Exemplars

The following illustrates the **stated** and **inferred** view for 738992009 |Chew (administration method)|:

![Diagram of exemplars](image)

### Dose Form Intended Site Hierarchy

#### Overview

The dose form intended site represents a general type of site of administration (e.g. cutaneous, nasal, oral, parenteral). To support fully defining concepts in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy, a hierarchy representing dose form intended site is required. Even though the concepts appear similar, dose form intended site should not be confused with route of administration, which is a concept used in dosage instructions for the administration of a particular medicinal product to a particular patient.

Concepts in the 736479009 |Dose form intended site (intended site)| hierarchy will be used to model concepts in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy; they will not be used to model concepts in the 763158003 |Medicinal product (product)| hierarchy.

The 736479009 |Dose form intended site (intended site)| hierarchy is a descendant of 362981000 |Qualifier value (qualifier value)|.

#### Modeling (stated view)

Descendants shall be modeled as follows.

| Parent concept | 736479009 |Dose form intended site (intended site) or a descendant of 736479009 |Dose form intended site (intended site) |
|----------------|----------------|
| Semantic tag   | (intended site) |

Exceptions: none
Definition status

900000000000074008|Necessary but not sufficient concept definition status (core metadata concept)

- Exceptions: none

Attributes

None

Naming Guidelines

FSN

Use the following pattern for the FSN where X is the intended site:

X (intended site)

Example:

- Oral (intended site)
- Otic (intended site)
- Parenteral (intended site)
- Vaginal (intended site)

Preferred Term

Use the following pattern for the PT where X is the intended site:

X

Example:

- Oral
- Otic
- Parenteral
- Vaginal

Synonyms

Synonyms are not allowed unless explicitly identified as an exception in the Editorial Guidelines.

Exceptions: none

Text Definitions

Required.

Exemplars

The following illustrates the stated and inferred view for 738984000 |Parenteral (intended site)|:

Dose Form Release Characteristic Hierarchy

Overview

The dose form release characteristic represents a general type of pattern of release (e.g. conventional, modified, delayed, prolonged) of the active ingredient substance(s) from the dose form. To support fully defining concepts in
the 736542009 |Pharmaceutical dose form (dose form)| hierarchy, a hierarchy representing dose form release characteristics is required.

Concepts in the 736480007 |Dose form release characteristic (release characteristic)| hierarchy will be used to model concepts in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy; they will not be used to model concepts in the 763158003 |Medicinal product (product)| hierarchy.

The 736480007 |Dose form release characteristic (release characteristic)| hierarchy is a descendant of 362981000| Qualifier value (qualifier value).

Modeling (stated view)

Descendants shall be modeled as follows:

| Parent concept | 736480007|Dose form release characteristic (release characteristic) or one or more descendants of 736480007|Dose form release characteristic (release characteristic) |
|----------------|-------------------------------------------------------------------------------------------------|
| Exceptions     | none                                                                                           |

| Semantic tag   | (release characteristic)                                                                           |
| Definition status | 90000000000000074008|Necessary but not sufficient concept definition status (core metadata concept) |
| Exceptions     | none                                                                                           |

| Attributes | None                                                                                           |

Naming Guidelines

<table>
<thead>
<tr>
<th>FSN</th>
<th>Use the following pattern for the FSN where X is the release characteristic, adding a hyphen when appropriate Combined release-characteristic concepts may be created to support modeling of concepts that display multiple release characteristics in a single formulation.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• X (release characteristic)</td>
</tr>
<tr>
<td></td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td>• Conventional release (release characteristic)</td>
</tr>
<tr>
<td></td>
<td>• Delayed-release (release characteristic)</td>
</tr>
<tr>
<td></td>
<td>• Prolonged-release (release characteristic)*</td>
</tr>
<tr>
<td></td>
<td>• Modified-release (release characteristic)**</td>
</tr>
<tr>
<td></td>
<td>• Delayed-release and prolonged-release (release characteristic)</td>
</tr>
<tr>
<td></td>
<td>*Prolonged was selected rather than Extended because feedback was received that it is more clear to translate for non-English speaking users; existing national drug extensions appear to use one or the other of these terms.</td>
</tr>
<tr>
<td></td>
<td>**Modified-release is a grouper that is not allowed to be used to model a Clinical drug concept.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Use the following pattern for the PT where X is the release characteristic, adding a hyphen when appropriate:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• X</td>
</tr>
<tr>
<td></td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td>• Conventional release</td>
</tr>
<tr>
<td></td>
<td>• Delayed-release</td>
</tr>
<tr>
<td></td>
<td>• Prolonged-release</td>
</tr>
<tr>
<td></td>
<td>• Modified-release</td>
</tr>
<tr>
<td></td>
<td>• Delayed-release and prolonged-release</td>
</tr>
</tbody>
</table>
### Exemplars

The following illustrates the **stated** and **inferred** view for 736849007 |Conventional release (release characteristic)|:

![Diagram 1](image1)

The following illustrates the **stated** and **inferred** view for 736848004 |Delayed-release (release characteristic)|:

![Diagram 2](image2)

The following illustrates the **stated** and **inferred** view for 737594003 |Delayed-release and prolonged-release (release characteristic)|:

![Diagram 3](image3)

### Dose Form Transformation Hierarchy

#### Overview

The dose form transformation represents a process where a dose form is transformed from that supplied by the manufacturer into a new dose form, usually to make it suitable for administration (e.g. dissolving a "powder for solution for injection" dose form into a "solution for injection" dose form). This may occur as part of the dispensing act or immediately before administration. To support fully defining concepts in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy, a hierarchy representing dose form transformation is required.
Concepts in the 736477006 |Dose form transformation (transformation)| hierarchy will be used to model concepts in the 736542009 |Pharmaceutical dose form (dose form)| hierarchy; they will not be used to model concepts in the 763158003 |Medicinal product (product)| hierarchy.

The 736477006 |Dose form transformation (transformation)| hierarchy is a descendant of 362981000|Qualifier value (qualifier value).

Modeling (stated view)

Descendants shall be modeled as follows.

| Parent concept | 736477006 |Dose form transformation (transformation)| or a descendant of 736477006 | Dose form transformation (transformation)|
|----------------|---------------------------------|---------------------------------|
| Exceptions     | none                            |                                  |

| Semantic tag   | (transformation)                 |

<table>
<thead>
<tr>
<th>Definition status</th>
<th>900000000000074008</th>
<th>Necessary but not sufficient concept definition status (core metadata concept)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exceptions</td>
<td>none</td>
<td></td>
</tr>
</tbody>
</table>

| Attributes        | None            |                                                                       |

Naming Guidelines

<table>
<thead>
<tr>
<th>FSN</th>
<th>Use the following pattern for the FSN where X is the transformation:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• X (transformation)</td>
</tr>
<tr>
<td>Example</td>
<td>Dissolve (transformation)</td>
</tr>
<tr>
<td></td>
<td>Disperse (transformation)</td>
</tr>
<tr>
<td></td>
<td>No transformation (transformation)</td>
</tr>
<tr>
<td></td>
<td>Disperse or dissolve (transformation)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Use the following pattern for the PT where X is the transformation:</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• X</td>
</tr>
<tr>
<td>Example</td>
<td>Dissolve</td>
</tr>
<tr>
<td></td>
<td>Disperse</td>
</tr>
<tr>
<td></td>
<td>No transformation</td>
</tr>
<tr>
<td></td>
<td>Disperse or dissolve</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Synonyms</th>
<th>Synonyms are not allowed unless explicitly identified as an exception in the Editorial Guidelines.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exceptions</td>
<td>none</td>
</tr>
</tbody>
</table>

| Text Definitions  | Required.                                                                                     |

Exemplars

The following illustrates the **stated** and **inferred** view for 761954006 |No transformation (transformation)|:
State of Matter Hierarchy

Overview
The state of matter represents a physical state of matter. To support fully defining concepts in the 736542009 | Pharmaceutical dose form (dose form) | hierarchy, a hierarchy representing state of matter is required.

Concepts in the 736471007 | State of matter (state of matter) | hierarchy will be used to model concepts in the 736478001 | Basic dose form (basic dose form) | hierarchy; they will not be used to model concepts in the 736542009 | Pharmaceutical dose form (dose form) | or 763158003 | Medicinal product (product) | hierarchies.

The 736471007 | State of matter (state of matter) | hierarchy is a descendant of 362981000 | Qualifier value (qualifier value) |

Modeling (stated view)
Descendants shall be modeled as follows.

| Parent concept | 736471007 | State of matter (state of matter) |
|----------------|-----------------------------|
| Exceptions: none |

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(state of matter)</th>
</tr>
</thead>
</table>
Definition status
900000000000074008|Necessary but not sufficient concept definition status (core metadata concept)
- Exceptions: none

Attributes
None

Naming Guidelines

FSN
Use the following pattern for the FSN where X is the state of matter:

X (state of matter)

Example:
- Gas (state of matter)
- Liquid (state of matter)
- Semi-solid (state of matter)
- Solid (state of matter)

Preferred Term
Use the following pattern for the PT where X is the state of matter:

X

Example:
- Gas
- Liquid
- Semi-solid
- Solid

Synonyms
Synonyms are not allowed unless explicitly identified as an exception in the Editorial Guidelines.
- Exceptions: none

Text Definitions
Required.

Exemplars
The following illustrates the stated and inferred view for 736679003 |Liquid (state of matter)|:

Appendix A - Cumulative Summary of Changes for Pharmaceutical Dose Form Editorial Guide

<table>
<thead>
<tr>
<th>Date</th>
<th>Comments</th>
<th>Effective date</th>
</tr>
</thead>
</table>

© Copyright 2021 International Health Terminology Standards Development Organisation
| 2021-Jan | Revisions to support making Pharmaceutical Dose Form SNOMED CT Editorial Guide available via SNOMED International Document Library. | 2021-Jan |
SNOMED CT Editorial Guide - Pharmaceutical and Biologic Product

This section provides editorial guidance for the 373873005 |Pharmaceutical / biologic product (product)| hierarchy and the supporting subhierarchy of 766940004 |Role (role)|

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Summary of Changes

<table>
<thead>
<tr>
<th>Date</th>
<th>Comments</th>
<th>Effective date</th>
</tr>
</thead>
<tbody>
<tr>
<td>2021-Jul</td>
<td>Updated to reflect:</td>
<td>2022-Jan</td>
</tr>
<tr>
<td></td>
<td>• Clinical drug concepts should not be modeling using &quot;lyophilized powder&quot;</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Pharmaceutical dose forms</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Clinical drug concepts with semi-solid dose form should have strength denominator normalized to /1 gram or /1 milliliter</td>
<td></td>
</tr>
</tbody>
</table>

Blood Product

410652009 |Blood product (product)| and its descendants will be retained as a primitive subhierarchy until use cases and/or detailed requirements are known.

Requests for addition of new concepts or modification of existing concepts will be evaluated on a case-by-case basis.

Bone Cements

356497001 |Bone cements (product)| and its descendants will be retained as a primitive subhierarchy until use cases and/or detailed requirements are known.

Requests for addition of new concepts or modification of existing concepts will be evaluated on a case-by-case basis.

Bone Graft Material

409248003 |Bone graft material (product)| and its descendants will be retained as a primitive subhierarchy until use cases and/or detailed requirements are known.

Requests for addition of new concepts or modification of existing concepts will be evaluated on a case-by-case basis.

Bone Morphogenic Protein Graft

411976009 |Bone morphogenic protein graft (product)| and its descendants will be retained as a primitive subhierarchy until use cases and/or detailed requirements are known.

Requests for addition of new concepts or modification of existing concepts will be evaluated on a case-by-case basis.

Dialysis Fluid

116178008 |Dialysis fluid (product)| and its descendants will be retained as a primitive subhierarchy until use cases and/or detailed requirements are known.
Requests for addition of new concepts or modification of existing concepts will be evaluated on a case-by-case basis.

**Dietary Product**

Development of editorial guidelines and related content updates for existing concept 373783004 |Dietary product (product)| and its descendants will be managed by the Nutrition Group as a future project. In the meantime, this hierarchy will be retained as a primitive subhierarchy.

Requests for addition of new concepts or modification of existing concepts will be evaluated on a case-by-case basis.

**Drug-Device Combination Product**

Development of editorial guidelines and related content updates for existing concept 411115002 |Drug-device combination product (product)| and its descendants will be done in conjunction with the Device Project. In the meantime, this hierarchy will be retained "as is".

Requests for addition of new concepts or modification of existing concepts will be evaluated on a case-by-case basis.

**Herbal Medicine**

349365008 |Herbal medicine (product)| and its descendants will be retained "as is" until use cases and/or detailed requirements are known.

Requests for addition of new concepts will be rejected. Requests for modification of existing concepts will be evaluated on a case-by-case basis.

**Homeopathic Medicine**

349363001 |Homeopathic medicine (product)| and its descendants will be retained "as is" until use cases and/or detailed requirements are known.

Requests for addition of new concepts will be rejected. Requests for modification of existing concepts will be evaluated on a case-by-case basis.

**Medicinal Product**

The grouper 763158003 |Medicinal product (product)|, a stated descendant of |Pharmaceutical / biologic product (product)|, was created to support top-level hierarchy changes in the future but avoids removing, renaming, or repurposing the existing |Pharmaceutical / biologic product (product)| concept.

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**Glossary for Medicinal Product**

The following definitions and abbreviations apply to this document:

<table>
<thead>
<tr>
<th>Term</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>1 Active immunity</td>
<td>The usually long lasting immunity which results from the production of antibodies by the immune system within an organism in response to the presence of an antigen.</td>
</tr>
<tr>
<td></td>
<td></td>
</tr>
<tr>
<td>---</td>
<td>---</td>
</tr>
<tr>
<td>2</td>
<td><strong>Active moiety</strong></td>
</tr>
<tr>
<td>3</td>
<td><strong>Adjuvant</strong></td>
</tr>
<tr>
<td>4</td>
<td><strong>Adsorption</strong></td>
</tr>
<tr>
<td>5</td>
<td><strong>Antibody</strong></td>
</tr>
<tr>
<td>6</td>
<td><strong>Antigen</strong></td>
</tr>
<tr>
<td>7</td>
<td><strong>BAN</strong></td>
</tr>
<tr>
<td>8</td>
<td><strong>Basis of Strength Substance</strong></td>
</tr>
<tr>
<td>9</td>
<td><strong>BoSS</strong></td>
</tr>
<tr>
<td>10</td>
<td><strong>CD</strong></td>
</tr>
<tr>
<td>11</td>
<td><strong>CDC</strong></td>
</tr>
</tbody>
</table>
| 12 | **Composite product** | Product that contains more than one single or multiple ingredient product packaged together. Example:  
  • PREVPAC® consists of a daily administration pack containing lansoprazole 30 mg oral capsules, amoxicillin 500 mg oral capsules, and clarithromycin 500 mg oral tablets. |
| 13 | **Conjugate vaccine** | Conjugate vaccines combine a weak antigen with a strong antigen (usually a protein/peptide carrier) so that the immune system has a stronger response to the weak antigen.  
  Conjugation is usually used for polysaccharide antigens, because polysaccharide antigens on their own produce only a B cell response (they are not whole cells, just pieces of pathogen cell wall). The conjugated peptide stimulate T cells which gives a more vigorous immune response and also promotes a more rapid and long-lasting immunologic memory (e.g Haemophilus influenzae type b conjugate vaccine, meningococcal conjugate vaccine). The carrier protein may be the diphtheria toxoid or the tetanus toxoid. |
<p>| | | |</p>
<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
</table>
| 14 | CVX code | The "vaccine administered" code set developed and maintained by the CDC's National Center of Immunization and Respiratory Diseases. When paired with a MVX (manufacturer) code, the specific trade named vaccine may be indicated. Each code is associated with a status indicating its availability in the United States (e.g. Active, Inactive, Non-US).
|   |   | [https://www2a.cdc.gov/vaccines/iis/iisstandards/vaccines.asp?rpt=cvx](https://www2a.cdc.gov/vaccines/iis/iisstandards/vaccines.asp?rpt=cvx) |
| 15 | Disposition | A “behavior” that an active ingredient will exhibit or participate in, given the appropriate context.
|   |   | Example:
|   |   | • 734727006 [Opioid receptor agonist (disposition)]
|   |   | • 734698003 [Beta adrenergic receptor antagonist (disposition)] |
| 16 | EHR | Electronic Health Record |
| 17 | FHIR | Fast Healthcare Interoperability Resources
|   |   | [https://www.hl7.org/fhir/](https://www.hl7.org/fhir/) |
| 18 | FSN | Fully Specified Name |
| 19 | Grouper | A concept designed to aggregate concepts based on specific characteristics or commonalities.
|   |   | Types of groupers included in the [Pharmaceutical / biologic product (product)] hierarchy are described in further detail in following sections of this document. |
| 20 | GTIN | Global Trade Item Number |
| 21 | Hapten | A molecule that is incapable, alone, of causing the production of antibodies but, can, however, combine with a larger antigenic molecule called a carrier to form an antigenic complex (see hapten-carrier complex). [Stedman's Medical Dictionary, adapted] |
| 22 | Hapten-carrier complex | An association between a hapten molecule and an antigen molecule that can stimulate production of antibodies, some of which combine with the hapten portion of the complex. [Stedman's Medical Dictionary, adapted] |
| 23 | Herbal medicine product | Herbal medicines include herbs, herbal materials, herbal preparations and finished herbal products, that contain as active ingredients parts of plants, or other plant materials, or combinations.
| 24 | HL7 | Health Level Seven International
| 25 | Homeopathic product | An alternative approach to medicine based on the belief that natural substances, prepared in a special way and used most often in very small amounts, restore health. According to these beliefs, in order for a remedy to be effective, it must cause in a healthy person the same symptoms being treated in the patient.
<p>|   |   | <a href="https://www.nhs.uk/conditions/homeopathy/#what-is-homeopathy">https://www.nhs.uk/conditions/homeopathy/#what-is-homeopathy</a> |</p>
<table>
<thead>
<tr>
<th>Code</th>
<th>Term</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>26</td>
<td>IDMP</td>
<td>Identification of Medicinal Products</td>
</tr>
<tr>
<td></td>
<td></td>
<td>A set of five standards developed by the International Organization for Standardization (ISO) for the identification of medicinal products.</td>
</tr>
<tr>
<td>27</td>
<td>IHTSDO</td>
<td>International Health Terminology Standards Development Organisation</td>
</tr>
<tr>
<td></td>
<td></td>
<td><a href="https://www.snomed.org/">https://www.snomed.org/</a></td>
</tr>
<tr>
<td></td>
<td></td>
<td>The trading name for IHTSDO is SNOMED International.</td>
</tr>
<tr>
<td>28</td>
<td>Immunogen</td>
<td>A complete antigen (i.e. can evoke the production of antibodies). Synonym for antigen...except that it is sometimes used without the specificity of the serotype (e.g. no statement of valency) whereas an antigen should have the valency specified.</td>
</tr>
<tr>
<td>29</td>
<td>Immunoglobulin</td>
<td>A class of polypeptide chain proteins in two pairs (one light, one heavy); antibodies are immunoglobulins and most immunoglobulins function as antibodies. The class of immunoglobulins also includes pathological proteins such as Bence Jones or myeloma globulins.</td>
</tr>
<tr>
<td>30</td>
<td>Inactivated vaccine product</td>
<td>A vaccine product whose antigenic content consists of the disease-causing pathogen that has been inactivated (&quot;killed&quot;) usually by heat or by chemicals such as formaldehyde. The pathogen cannot replicate itself at all, but it is still intact and can therefore evoke antibody production (example: polio vaccine).</td>
</tr>
<tr>
<td>31</td>
<td>INN</td>
<td>International Nonproprietary Name</td>
</tr>
<tr>
<td>32</td>
<td>International Nonproprietary Name</td>
<td>INNs facilitate the identification of pharmaceutical substances or active pharmaceutical ingredients. Each INN is a unique name that is globally recognized and is public property. A nonproprietary name is also known as a generic name. For more information: <a href="http://www.who.int/medicines/services/inn/en/">http://www.who.int/medicines/services/inn/en/</a> To search for INNs: <a href="https://mednet-communities.net/inn/db/searchinn.aspx">https://mednet-communities.net/inn/db/searchinn.aspx</a></td>
</tr>
<tr>
<td>33</td>
<td>ISO</td>
<td>International Organization for Standardization</td>
</tr>
<tr>
<td></td>
<td></td>
<td><a href="https://www.iso.org/home.html">https://www.iso.org/home.html</a></td>
</tr>
<tr>
<td>34</td>
<td>Live attenuated vaccine product</td>
<td>A vaccine product whose antigenic content is derived from the disease-causing pathogen but which has been altered to make it less virulent. The pathogen in a live attenuated vaccine has lost its ability to replicate in human cells but still viable to evoke antibody production (e.g. measles, mumps, and rubella vaccine, varicella vaccine).</td>
</tr>
<tr>
<td>35</td>
<td>Monovalent vaccine</td>
<td>A vaccine product that contains a single antigenic serotype.</td>
</tr>
<tr>
<td>36</td>
<td>MP</td>
<td>Medicinal Product</td>
</tr>
<tr>
<td>37</td>
<td>MPF</td>
<td>Medicinal Product Form</td>
</tr>
</tbody>
</table>
| 38   | Multiple ingredient product | Product that contains more than one active ingredient in a single manufactured dose form. Example:  
- 377265005 |Product containing precisely captopril 50 milligram and hydrochlorothiazide 15 milligram/1 each conventional release oral tablet [clinical drug]  
- 407853009 |Product containing precisely codeine phosphate 15 milligram and paracetamol 500 milligram/1 each conventional release oral tablet [clinical drug] |
<table>
<thead>
<tr>
<th>No.</th>
<th>Term</th>
<th>Definition</th>
</tr>
</thead>
<tbody>
<tr>
<td>39</td>
<td>MVX Code</td>
<td>The &quot;Manufacturers of vaccines&quot; code set developed and maintained by the CDC’s National Center of Immunization and Respiratory Diseases. When paired with a CVX (vaccine administered) code, the specific trade name defined vaccine may be indicated. Each code is associated with a status indicating if the manufacturer is currently making vaccines for distribution in the United States (e.g. Active, Inactive).</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td><a href="https://www2a.cdc.gov/vaccines/iis/iisstandards/vaccines.asp?rpt=mvx">https://www2a.cdc.gov/vaccines/iis/iisstandards/vaccines.asp?rpt=mvx</a></td>
</tr>
<tr>
<td>40</td>
<td>Passive immunity</td>
<td>The time limited, usually short lived, immunity acquired by direct transference of antibodies into an organism (e.g. by injection of immunoglobulin).</td>
</tr>
<tr>
<td>41</td>
<td>Polyclonal vaccine</td>
<td>A vaccine product that contains multiple antigenic serotypes; the number of which may be stated (e.g. tetravalent (4), pentavalent (5)).</td>
</tr>
<tr>
<td>42</td>
<td>Precise active ingredient</td>
<td>The actual active ingredient that is contained in the product.</td>
</tr>
<tr>
<td>43</td>
<td>Proximal primitive modeling pattern</td>
<td>The proximal primitive modeling pattern is the preferred SNOMED modeling approach.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Note: Additional information about proximal primitive modeling and intermediate primitive concepts can be found in the main SNOMED CT Editorial Guide.</td>
</tr>
<tr>
<td>44</td>
<td>PT</td>
<td>Preferred Term</td>
</tr>
<tr>
<td>45</td>
<td>Role</td>
<td>Role is associated with a particular purpose or outcome. The role is often by virtue of its manufacture or use. Since all occurrences of a given substance or drug product may not be used in the same manner roles are not always and necessarily true.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• 53009005</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• 27867009</td>
</tr>
<tr>
<td>46</td>
<td>Serotype</td>
<td>A subdivision of a species or subspecies distinguishable from other strains therein on the basis of antigenicity. [Stedman’s Medical Dictionary]</td>
</tr>
<tr>
<td>47</td>
<td>Single ingredient product</td>
<td>Product that contains one and only one active ingredient in a single manufactured dose form.</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• 765732008</td>
</tr>
<tr>
<td></td>
<td></td>
<td>• 446347007</td>
</tr>
<tr>
<td>48</td>
<td>Subunit vaccine</td>
<td>A vaccine product whose antigenic content is a target part of a pathogen (e.g. a specific protein from the pathogen) rather than the whole pathogen, produced either by isolation from the pathogen or by recombinant technology.</td>
</tr>
<tr>
<td>49</td>
<td>Toxoid</td>
<td>A vaccine product whose antigenic content is a toxin produced by a pathogen that has been treated, commonly with formaldehyde, so as to destroy its toxic property but retain its antigenicity (i.e. its capability of stimulating the production of antitoxin antibodies and thus of producing an active immunity).</td>
</tr>
<tr>
<td>50</td>
<td>Traditional medicine product</td>
<td>Traditional medicine is the sum total of the knowledge, skills, and practices based on the theories, beliefs, and experiences indigenous to different cultures, whether explicable or not, used in the maintenance of health as well as in the prevention, diagnosis, improvement or treatment of physical and mental illness.</td>
</tr>
</tbody>
</table>
|     |                                          | https://www.who.int/traditional-complementary-integrative-medicine/en/
<table>
<thead>
<tr>
<th></th>
<th></th>
<th>Unified Code for Units of Measure</th>
</tr>
</thead>
<tbody>
<tr>
<td>UCUM</td>
<td></td>
<td><a href="http://unitsofmeasure.org/trac">http://unitsofmeasure.org/trac</a></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th></th>
<th>United States Adopted Name</th>
</tr>
</thead>
<tbody>
<tr>
<td>USAN</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th></th>
<th>Vaccine</th>
</tr>
</thead>
</table>
|Vaccine | | Any preparation intended for active immunologic prophylaxis (e.g. preparation of killed microns of virulent strains or living microbes of attenuated (variant or mutant) strain; or microbial, fungal, plant, protozoal or metazoan derivatives or products. [Stedman’s Medical Dictionary] 

Originally only applied to live vaccine (vaccinia, cowpox) virus inoculated in the skin as prophylaxis against smallpox and obtained from the sin of calves inoculated with seed virus. |

<table>
<thead>
<tr>
<th></th>
<th></th>
<th>Vaccine pharmacovigilance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vaccine pharmacovigilance</td>
<td></td>
<td>Vaccine pharmacovigilance is defined by the WHO as &quot;the science and activities relating to the detection, assessment, understanding and communication of adverse events following immunization and other vaccine- or immunization-related issues, and to the prevention of untoward effects of the vaccine or immunization&quot;. <a href="https://www.who.int/vaccine_safety/initiative/tools/CIOMS_report_WG_vaccine.pdf">https://www.who.int/vaccine_safety/initiative/tools/CIOMS_report_WG_vaccine.pdf</a></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th></th>
<th>The number of antigenic serotypes present in a vaccine product.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vaccine valency</td>
<td>Antigenic valency</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th></th>
<th></th>
<th>World Health Organization</th>
</tr>
</thead>
<tbody>
<tr>
<td>WHO</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

### Assumptions, Requirements, and Scope for Medicinal Product

#### General Assumptions and Requirements

<table>
<thead>
<tr>
<th>Assumption or Requirement</th>
</tr>
</thead>
</table>
| 1 | Concept model will conform to description logic principles, including use of the classifier to organize the concepts in the hierarchy where appropriate. 
  
  - After classification, the top level concepts in the hierarchy will primarily be sufficiently defined grouper concepts. |
| 2 | Concept model will include attributes necessary to sufficiently define the concepts to ensure consistent and reproducible modeling of concepts. |
| 3 | Any requirement to align to external standards or registries will be explicitly documented. 
  
  Concept model will be compatible with IDMP standards (where appropriate). |
| 4 | Concepts shall be sufficiently defined using proximal primitive modeling methodology unless explicitly noted as an exception in the editorial guidelines. 
  
  - Note: Additional information about proximal primitive modeling and intermediate primitive concepts can be found in the main SNOMED CT Editorial Guide. |
| 5 | Concept model will not support universal restrictions. |
| 6 | Concept model will not support nesting. |
| 7 | Content in the [Medicinal product] hierarchy in the International Release is not intended to support prescribing use cases but may be sufficient to do so for some implementations. |
8. Content in the [Medicinal product] hierarchy in the International Release is not intended to eliminate the need for a national extension.

### Scope of Content

<table>
<thead>
<tr>
<th>Scope of Content</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Age ranges</td>
</tr>
<tr>
<td>• Concepts that refer to age ranges (e.g. adult, pediatric, infant, junior, adolescent) are out of scope.</td>
</tr>
<tr>
<td>• Exception: Vaccine products MP-only Concepts maybe modeled with Has target population attribute that specifies a target population.</td>
</tr>
<tr>
<td>2. Adjuvants</td>
</tr>
<tr>
<td>• Concepts that refer to adjuvants are out of scope.</td>
</tr>
<tr>
<td>3. Ayurvedics</td>
</tr>
<tr>
<td>• Concepts that refer to Ayurvedic medicine concepts are out of scope.</td>
</tr>
<tr>
<td>4. Brand names</td>
</tr>
<tr>
<td>• Concepts that refer to brand names are out of scope.</td>
</tr>
<tr>
<td>5. Colors</td>
</tr>
<tr>
<td>• Concepts that refer to color (e.g. color of tablet, capsule, or solution) are out of scope.</td>
</tr>
<tr>
<td>6. Composite products</td>
</tr>
<tr>
<td>• Concepts that refer to composite products are out of scope.</td>
</tr>
<tr>
<td>7. Excipients</td>
</tr>
<tr>
<td>• Concepts that refer to excipients are out of scope.</td>
</tr>
<tr>
<td>8. Flavors</td>
</tr>
<tr>
<td>• Concepts that refer to flavor are out of scope.</td>
</tr>
<tr>
<td>9. Investigational products</td>
</tr>
<tr>
<td>• Concepts that refer to products under development but not marketed in any member country are out of scope.</td>
</tr>
<tr>
<td>• Exceptions may be made on a case-by-case basis (e.g. adding investigational products that are being widely used in pandemic).</td>
</tr>
<tr>
<td>10. No longer marketed or available for sale</td>
</tr>
<tr>
<td>• Existing concepts representing products that are no longer marketed or available for sale will be retained as active concepts in the International Release.</td>
</tr>
<tr>
<td>• Requests for addition of new concepts representing products that are known to be no longer marketed or available for sale will be considered for inclusion on a case-by-case basis.</td>
</tr>
<tr>
<td>11. Non-human use</td>
</tr>
<tr>
<td>• Concepts that represent products intended only for non-human use are out of scope.</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>---</td>
</tr>
<tr>
<td>12</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>13</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>14</td>
</tr>
<tr>
<td>15</td>
</tr>
<tr>
<td></td>
</tr>
<tr>
<td>16</td>
</tr>
<tr>
<td></td>
</tr>
</tbody>
</table>

Naming and Modeling Guidelines for Medicinal Product

Groupers Based on Disposition

Overview

Grouper concepts based on disposition of an active ingredient that can be sufficiently defined may be included in the [763158003 |Medicinal product (product)] hierarchy.

Note: This section applies to grouper concepts representing a single disposition. Groupers comprised of multiple dispositions are described in a separate section.

A high-level grouper concept supports the organization of the medicinal product concepts based on disposition:

- 766779001 |Medicinal product categorized by disposition (product)|

Modeling (stated view)

<table>
<thead>
<tr>
<th>Stated parent concept</th>
<th>763158003</th>
<th>Medicinal product (product)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Semantic tag</td>
<td>(product)</td>
<td></td>
</tr>
<tr>
<td>Definition status</td>
<td>900000000000073002</td>
<td>Sufficiently defined by necessary conditions definition status (core metadata concept)</td>
</tr>
<tr>
<td>Attribute: Has active ingredient</td>
<td>Range: &lt;=105590001</td>
<td>Substance (substance)</td>
</tr>
<tr>
<td></td>
<td>• NOTE: While the allowed range is broader, the</td>
<td>Medicinal product</td>
</tr>
<tr>
<td></td>
<td>• Cardinality: 0..*</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• NOTE: While the allowed range is broader, the</td>
<td>Medicinal product</td>
</tr>
</tbody>
</table>
Naming Guidelines

**FSN**

Use the following pattern for the FSN; align naming and case sensitivity with the PT for the concept that is selected as the 726542003 |Has disposition (attribute)| attribute value for the substance concept used as the attribute value for the 127489000 |Has active ingredient (attribute)|:

- Product containing `<Active ingredient PT>` (product)

Example:

- Product containing histamine receptor antagonist (product)
- Product containing histamine H2 receptor antagonist (product)

**Preferred Term**

Use the following pattern for the PT; align naming and case significance with the PT for the concept that is selected as the 726542003 |Has disposition (attribute)| attribute value for the substance concept used as the attribute value for the 127489000 |Has active ingredient (attribute)|:

- `<Active ingredient PT>`-containing product

Example:

- Histamine receptor antagonist-containing product
- Histamine H2 receptor antagonist-containing product

**Synonyms**

Synonyms matching the FSN are not required.

Exemplars

The following illustrates the **stated** view for grouper concept 6425004 |Product containing histamine receptor antagonist (product)|:

![Diagram 1]

The following illustrates the **inferred** view for grouper concept 6425004 |Product containing histamine receptor antagonist (product)|:

![Diagram 2]

The following illustrates the **stated** view for grouper concept 108661002 |Product containing histamine H2 receptor antagonist (product)|:

![Diagram 3]
Groupers Based on Structure

Overview

Grouper concepts based on chemical structure of an active ingredient that can be sufficiently defined may be included in the [Medicinal product] hierarchy.

Note: This section applies to grouper concepts representing a single structure.

A high-level grouper concept supports the organization of the hierarchy based on structure:
- 763760008 [Medicinal product categorized by structure (product)]

Modeling (stated view)

Grouper concepts based on structure shall be modeled using the proximal primitive modeling pattern.

| Stated parent concept | 763158003|Medicinal product (product) |
|-----------------------|----------------------------------|
| Semantic tag          | (product)                        |
| Definition status     | 9000000000000073002|Sufficiently defined by necessary conditions definition status (core metadata concept)|
| Attribute: Has active ingredient | Range: &lt;105590001|Substance (substance) |
|                       | • NOTE: While the allowed range is broader, the [Medicinal product] grouper concepts based on structure should only use primitive grouper concepts that are descendants of 312413002 |Substance categorized by structure (substance)| as attribute values. |
|                       | • Cardinality: 0..* |
|                       | • NOTE: While the allowed range is broader, the [Medicinal product] grouper concepts based on structure should have one and only one |Has active ingredient| attribute. |
Naming Guidelines

<table>
<thead>
<tr>
<th>FSN</th>
<th>Use the following pattern for the FSN; align naming and case sensitivity with the PT for the concept that is selected as the attribute value for the 127489000 [Has active ingredient (attribute)], excluding the semantic tag.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Product containing <code>&lt;Active ingredient PT&gt;</code> (product)</td>
</tr>
<tr>
<td></td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td>• Product containing prostaglandin (product)</td>
</tr>
<tr>
<td></td>
<td>• Product containing A series prostaglandin (product)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Use the following pattern for the PT; align naming and case significance with the PT for the concept that is selected as the attribute value.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• <code>&lt;Active ingredient PT&gt;</code>-containing product</td>
</tr>
<tr>
<td></td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td>• Prostaglandin-containing product</td>
</tr>
<tr>
<td></td>
<td>• A series prostaglandin-containing product</td>
</tr>
</tbody>
</table>

| Synonyms | Synonyms matching the FSN are not required.                                                                                                                                                                                                               |

Exemplars

The following illustrates the **stated** view for 350067007 [Product containing prostaglandin (product)]:

```
350067007
Product containing prostaglandin (product)
```

```
703158003
Medicinal product (product)
```

```
127489000
Has active ingredient (attribute)
```

```
26351002
Prostaglandin (substance)
```

The following illustrates the **inferred** view for 350067007 [Product containing prostaglandin (product)]:

```
350067007
Product containing prostaglandin (product)
```

```
768626007
Product containing carboxylic acid and carboxylic acid derivative (product)
```

```
127489000
Has active ingredient (attribute)
```

```
26351002
Prostaglandin (substance)
```

The following illustrates the **stated** view for 350068002 [Product containing A series prostaglandin (product)]:

```
Groupers Based on Multiple Dispositions and/or Structures

Overview

Groupers comprised of two or more existing disposition and/or structure groupers that can be sufficiently defined may be included in the Medicinal product hierarchy.

High-level grouper concepts support the organization of the combined groupers based on disposition and/or structure:

- 766779001 |Medicinal product categorized by disposition (product)|
- 763760008 |Medicinal product categorized by structure (product)|

Modeling (stated view)

Combined grouper concepts shall be modeled using the proximal primitive modeling pattern.

| Stated parent concept | 763158003|Medicinal product (product) |
|-----------------------|---------------------------------|
| Semantic tag          | (product)                         |
| Definition status     | 9000000000000073002 |Sufficiently defined by necessary conditions definition status (core metadata concept)|
### Attribute: Has active ingredient

- **Range:** `<<105590001|Substance (substance)`
  - **NOTE:** While the allowed range is broader, the `|Medicinal product|` combined grouper concepts based on disposition and/or structure should only use sufficiently defined grouper concepts that are descendants of `766739005 |Substance categorized by disposition (substance)|` and/or primitive grouper concepts that are descendants of `312413002 |Substance categorized by structure (substance)|` as attribute values.

- **Cardinality:** `0..*`
  - **NOTE:** While the allowed range is broader, the `|Medicinal product|` combined grouper concepts should have one or more `|Has active ingredient|` attributes.

### Naming Guidelines

#### FSN

- Use the following pattern for the FSN if the combined grouper is comprised of two dispositions or two structural groupers; align naming and case significance with the PT as described in Section 4.1 and 4.2, respectively. The active ingredients must be in alphabetical order and separated by the word “and”.

  - Product containing `<Active ingredient PT>` and `<Active ingredient PT>` (product)

  **Example:**

  - Product containing norepinephrine reuptake inhibitor and serotonin reuptake inhibitor (product)

- Use the following pattern for the FSN if the combined grouper is comprised of one disposition and one structural grouper; align naming and case significance with the FSN for the concept with the FSN as described in Section 4.1 and 4.2, respectively.

  - Product containing `<Structural grouper active ingredient PT>` and `<Disposition grouper active ingredient PT>` (product)

  **Example:**

  - Product containing piperazine derivative and histamine receptor antagonist (product)
Preferred Term

Use the following pattern for the PT if the combined grouper is comprised of two dispositions or two structural groupers; align naming and case significance with the PT for the concept that is selected as the attribute value. The active ingredients must be in alphabetical order and separated by the word “and”.

- `<Active ingredient PT>` and `<Active ingredient PT>`-containing product

Example:

- Norepinephrine reuptake inhibitor and serotonin reuptake inhibitor-containing product

Use the following pattern for the FSN if the combined grouper is comprised of one disposition and one structural grouper; align naming and case significance with the FSN for the concept that is selected as the attribute value.

- `<Structural grouper active ingredient PT>` and `<Disposition grouper active ingredient PT>`-containing product

Example:

- Piperazine derivative and histamine receptor antagonist-containing product

Synonyms

Synonyms matching the FSN are not required.

Exemplars

The following illustrates the stated view for combined grouper concept 767562003 |Product containing norepinephrine reuptake inhibitor and serotonin reuptake inhibitor (product)|:

The following illustrates the inferred view for combined grouper concept 767562003 |Product containing norepinephrine reuptake inhibitor and serotonin reuptake inhibitor (product)|:

The following illustrates the stated view for combined grouper concept 70343008 |Product containing piperazine derivative and histamine receptor antagonist (product)|:
The following illustrates the inferred view for combined grouper concept 70343008 |Product containing piperazine derivative and histamine receptor antagonist (product)|:

Groupers Based on Dose form intended site

Overview

Groupers based on "Dose form intended site" that can be sufficiently defined may be included in the 763158003|Medicinal product (product)| hierarchy.

Example:
- Product manufactured as oral dose form (product)
- Product manufactured as parenteral dose form (product)

Modeling (stated view)

Grouper concepts based on "Dose form intended site" shall be modeled using the proximal primitive modeling pattern.

| Stated parent                  | 763158003|Medicinal product (product) |
|--------------------------------|------------------------------------------------|
| Semantic tag                  | (product)                                      |
| Definition status             | 90000000000000073002 |Sufficiently defined by necessary conditions definition status (core metadata concept)|
| Attribute:                    | Has manufactured dose form                      |
|                               | • Range: <<736542009|Pharmaceutical dose form (dose form) |
|                               |     • NOTE: While the allowed range is broader, the |Medicinal product| grouper concepts based on dose form intended site should use sufficiently defined grouper concepts that are descendants of 736542009|Pharmaceutical dose form (dose form)| representing intended site as attribute values. |
|                               |     • Cardinality: 0..*                          |
|                               |     • NOTE: While the allowed range is broader, the |Medicinal product| grouper concepts based on disposition should have one and only one |Has manufactured dose form| attribute. |
Naming Guidelines

<table>
<thead>
<tr>
<th>FSN</th>
<th>Use the following pattern for the FSN; align naming and case significance with the FSN for the concepts that are selected as the attribute value, excluding the semantic tag.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Product manufactured as <code>&lt;Manufactured dose form FSN&gt; (product)</code> Example:</td>
</tr>
<tr>
<td></td>
<td>• Product manufactured as oral dose form (product)</td>
</tr>
<tr>
<td></td>
<td>• Product manufactured as parenteral dose form (product)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Use the following pattern for the Preferred Term; align naming and case significance with the PT for the concept that is selected as the attribute value.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Product manufactured as <code>&lt;Manufactured dose form PT&gt;</code> Example:</td>
</tr>
<tr>
<td></td>
<td>• Product manufactured as oral dose form</td>
</tr>
<tr>
<td></td>
<td>• Product manufactured as parenteral dose form</td>
</tr>
</tbody>
</table>

| Synonyms | Synonyms are not generally created.                                                                                         |

Exemplars

The following illustrates the stated and inferred view for 440131009 |Product manufactured as oral dose form (product)|:

The following illustrates the stated and inferred view for 440132002 |Product manufactured as parenteral dose form (product)|:
Medicinal Product containing Concept

Overview
The Medicinal Product "containing" (MP-containing) concept is an abstract representation of the active ingredient(s) for a medicinal product. It means that the medicinal product must contain the active ingredient(s) specified in the FSN but may also contain a modification of the active ingredient(s) specified in the FSN or may contain additional active ingredient(s).

Example:
- Product containing axitinib (medicinal product)
- Product containing abacavir and lamivudine (medicinal product)

Modeling (stated view)
MP-containing concepts shall be modeled using the proximal primitive modeling pattern.

<table>
<thead>
<tr>
<th>Stated parent concept</th>
<th>763158003</th>
<th>Medicinal product (product)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Semantic tag</td>
<td>(medicinal product)</td>
<td></td>
</tr>
<tr>
<td>Definition status</td>
<td>900000000000073002</td>
<td>Sufficiently defined by necessary conditions definition status (core metadata concept)</td>
</tr>
<tr>
<td>Attribute:</td>
<td>Has active ingredient</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Range: 105590001</td>
<td>Substance (substance) - descendants only, excluding concepts representing structural groupers, dispositions, or combined substances</td>
</tr>
<tr>
<td></td>
<td>• Cardinality: 1..* - there is no technical limit on the number of Has active ingredient attributes that may be added to a concept; a practical limit may be imposed at a later date.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>For content in the International Release, this attribute value should represent the base ingredient, not a modification, unless explicitly identified as an exception.</td>
<td></td>
</tr>
<tr>
<td></td>
<td>• Exceptions:</td>
<td></td>
</tr>
<tr>
<td></td>
<td>▪ Chemical element with multiple modification (e.g. 422232005</td>
<td>Calcium lactate gluconate (substance)</td>
</tr>
<tr>
<td></td>
<td>▪ Chloral hydrate (e.g. 28748001</td>
<td>Product containing chloral hydrate (medicinal product)</td>
</tr>
<tr>
<td></td>
<td>▪ Liposome or lipid complex substances (e.g. 426490000</td>
<td>Vincristine liposome (substance)</td>
</tr>
<tr>
<td></td>
<td>▪ Pegylated substance (e.g. 3855440005</td>
<td>Pegfilgrastim (substance)</td>
</tr>
<tr>
<td></td>
<td>▪ Radiopharmaceutical (e.g. 783864004</td>
<td>Product containing cyanocobalamin (58-Co) (medicinal product)</td>
</tr>
<tr>
<td></td>
<td>▪ Silver sulfadiazine (e.g. 19630009</td>
<td>Product containing silver sulfadiazine (medicinal product)</td>
</tr>
</tbody>
</table>
Naming Guidelines

| FSN | Use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concept that is selected as the attribute value, excluding the semantic tag. For multiple ingredient drug products, the active ingredients must be in alphabetical order and separated by the word “and”.
<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>• Product containing <code>&lt;Active ingredient FSN&gt;</code> (medicinal product)</td>
<td></td>
</tr>
<tr>
<td>• Product containing <code>&lt;Active ingredient FSN&gt;</code> and <code>&lt;Active ingredient FSN&gt;</code> (medicinal product)</td>
<td></td>
</tr>
<tr>
<td>• Product containing <code>&lt;Active ingredient FSN&gt;</code> and <code>&lt;Active ingredient FSN&gt;</code> and <code>&lt;Active ingredient FSN&gt;</code> (medicinal product)</td>
<td></td>
</tr>
<tr>
<td>Example:</td>
<td></td>
</tr>
<tr>
<td>• Product containing axitinib (medicinal product)</td>
<td></td>
</tr>
<tr>
<td>• Product containing abacavir and lamivudine (medicinal product)</td>
<td></td>
</tr>
<tr>
<td>• Product containing abacavir and lamivudine and zidovudine (medicinal product)</td>
<td></td>
</tr>
</tbody>
</table>

| Preferred Term | Use the following pattern for the PT; align naming and case sensitivity with the PT for the concept that is selected as the attribute value. For multiple ingredient drug products, the active ingredients must be in alphabetical order and separated by the word “and”.
<table>
<thead>
<tr>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>• <code>&lt;Active ingredient PT&gt;</code>-containing product</td>
<td></td>
</tr>
<tr>
<td>• <code>&lt;Active ingredient PT&gt;</code>- and <code>&lt;Active ingredient PT&gt;</code>-containing product</td>
<td></td>
</tr>
<tr>
<td>• <code>&lt;Active ingredient PT&gt;</code>- and <code>&lt;Active ingredient PT&gt;</code>- and <code>&lt;Active ingredient PT&gt;</code>-containing product</td>
<td></td>
</tr>
<tr>
<td>Example:</td>
<td></td>
</tr>
<tr>
<td>• Axitinib-containing product</td>
<td></td>
</tr>
<tr>
<td>• Abacavir- and lamivudine-containing product</td>
<td></td>
</tr>
<tr>
<td>• Abacavir- and lamivudine- and zidovudine-containing product</td>
<td></td>
</tr>
</tbody>
</table>

| Synonym | Synonyms matching the FSN are not required. |

Exemplars

The following illustrates the **stated** view for 714627007 |Product containing aflibercept (medicinal product)|:

![Diagram](image)

The following illustrates the **inferred** view for 714627007 |Product containing aflibercept (medicinal product)|:
The following illustrates the **stated** view for 412556009 |Product containing codeine and paracetamol (medicinal product)|:

![Stated view diagram](image)

The following illustrates the **inferred** view for 412556009 |Product containing codeine and paracetamol (medicinal product)|:

![Inferred view diagram](image)
Medicinal Product containing only Concept

Overview
The Medicinal Product “containing only” (MP-only) concept is an abstract representation of the active ingredient(s) for a medicinal product. It means that the medicinal product must contain only the active ingredient(s) specified in the FSN but may also contain a modification of the active ingredient(s) specified in the FSN.

Example:
- Product containing only axitinib (medicinal product)
- Product containing only abacavir and lamivudine (medicinal product)

Modeling (stated view)
MP-only concepts shall be modeled using the proximal primitive modeling pattern.

| Stated parent concept | 763158003|Medicinal product (product) |
|-----------------------|----------------|
| Semantic tag          | (medicinal product) |
| Definition status     | 90000000000073002 [Sufficiently defined by necessary conditions definition status (core metadata concept)] |
### Attribute: Has active ingredient

- **Range:** 105590001|Substance (substance) - descendants only, excluding concepts representing structural groupers, dispositions, or combined substances
- **Cardinality:** 1..* - there is no technical limit on the number of Has active ingredient attributes that may be added to a concept; a practical limit may be imposed at a later date.

For content in the International Release, this attribute value should represent the base ingredient, not a modification, unless explicitly identified as an exception.

- **Exceptions:**
  - Chemical element with multiple modification (e.g. 422232005 |Calcium lactate gluconate (substance)|)
  - Chloral hydrate (e.g. 775158004 |Product containing only chloral hydrate (medicinal product)|)
  - Liposome or lipid complex substances (e.g. 426490000 |Vincristine liposome (substance)|, 425953004 |Amphotericin B lipid complex (substance)|, 768664009 |Amphotericin B phospholipid complex (substance)|, 427544000 |Amphotericin B cholesteryl sulfate complex (substance)|)
  - Pegylated substance (e.g. 385544005 |Pegfilgrastim (substance)|, 770965008 |Pegvaliase (substance)|)
  - Radiopharmaceutical (e.g. 783865003 |Product containing only cyanocobalamin (58-Co) (medicinal product)|, 783855009 |Product containing only sodium iodide (131-I) (medicinal product)|)
  - Silver sulfadiazine (e.g. 864008004 |Product containing only silver sulfadiazine (medicinal product)|)

### Attribute: Count of base of active ingredient

- **Concrete Type:** Integer
- **Range:** >0..1
- **Cardinality:** 1..1

### Naming Guidelines

Use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concept that is selected as the attribute value, excluding the semantic tag. For multiple ingredient drug products, the active ingredients must be in alphabetical order and separated by the word “and”.

- Product containing only `<Active ingredient FSN>` (medicinal product)
- Product containing only `<Active ingredient FSN>` and `<Active ingredient FSN>` (medicinal product)
- Product containing only `<Active ingredient FSN>` and `<Active ingredient FSN>` and `<Active ingredient FSN>` (medicinal product)

**Example:**

- Product containing only axitinib (medicinal product)
- Product containing only abacavir and lamivudine (medicinal product)
- Product containing only abacavir and lamivudine and zidovudine (medicinal product)
Use the following pattern for the PT; align naming and case sensitivity with the PT for the concept that is selected as the attribute value. For multiple ingredient drug products, the active ingredients must be in alphabetical order and separated by the word "and".

- `<Active ingredient PT>` only product
- `<Active ingredient PT>` and `<Active ingredient PT>` only product
- `<Active ingredient PT>` and `<Active ingredient PT>` and `<Active ingredient PT>` only product

Example:

- Axitinib only product
- Abacavir and lamivudine only product
- Abacavir and lamivudine and zidovudine only product

Exemplars

The following illustrates the **stated** view for 773390003 |Product containing only axitinib (medicinal product)|:

The following illustrates the **inferred** view for 773390003 |Product containing only axitinib (medicinal product)|:
The following illustrates the **stated** view for 775360007 |Product containing only codeine and paracetamol (medicinal product)|:

```
775360007
Product containing only codeine and paracetamol (medicinal product)
```

```
175738008
Medicinal product (product)
```

```
114213005
Count of base of active ingredient (attribute)
```

```
127490000
Has active ingredient (attribute)
```

```
387494007
Codeine (substance)
```

```
127490000
Has active ingredient (attribute)
```

```
387517004
Paracetamol (substance)
```

The following illustrates the **inferred** view for 775360007 |Product containing only codeine and paracetamol (medicinal product)|:

```
775360007
Product containing only codeine and paracetamol (medicinal product)
```

```
41566000
Product containing codeine and paracetamol (medicinal product)
```

```
114213005
Count of base of active ingredient (attribute)
```

```
127490000
Has active ingredient (attribute)
```

```
387494007
Codeine (substance)
```

```
127490000
Has active ingredient (attribute)
```

```
387517004
Paracetamol (substance)
```

**Medicinal Product Form containing Concept**

**Overview**

The Medicinal Product Form "containing" (MPF-containing) concept is an abstract representation of active ingredient(s) and dose form intended site for a medicinal product. It means that the medicinal product must contain the active ingredient(s) specified in the FSN but may also contain a modification of the active ingredient(s) specified in the FSN or may contain additional active ingredient(s) as well.

**Example:**

- Product containing axitinib in oral dose form (medicinal product form)
- Product containing abacavir and lamivudine in oral dose form (medicinal product form)
Modeling (stated view)

MPF-containing concepts shall be modeled using the proximal primitive modeling pattern.

| Stated parent concept | 763158003|Medicinal product (product) |
| Semantic tag | (medicinal product form) |
| Definition status | 900000000000073002 |Sufficiently defined by necessary conditions definition status (core metadata concept) |
| Attribute: | |
| Has active ingredient | • Range: 105590001|Substance (substance) - descendants only, excluding concepts representing structural groupers, dispositions, or roles |
| | • Cardinality: 1..* - there is no technical limit on the number of Has active ingredient attributes that may be added to a concept; a practical limit may be imposed at a later date. |
| For content in the International Release, this attribute value should represent the base ingredient, not a modification, unless explicitly identified as an exception. |
| • Exceptions: | |
| | • Chemical element with multiple modification (e.g. 422232005 |Calcium lactate gluconate (substance)) |
| | • Chloral hydrate [e.g. 386735001 |Product containing chloral hydrate in oral dose form (medicinal product form)] |
| | • Liposome or lipid complex substances(e.g. 426490000 |Vincristine liposome (substance)], 425953004 |Amphotericin B lipid complex (substance)], 768664009 |Amphotericin B phospholipid complex (substance)], 427544000 |Amphotericin B cholesteryl sulfate complex (substance)) |
| | • Pegylated substance (e.g. 385544005 |Pegfilgrastim (substance)], 770965008 |Pegvaliase (substance)) |
| | • Radiopharmaceutical (e.g. 783866002 |Product containing cyanocobalamin (58-Co) in oral dose form (medicinal product form)], 783856005 |Product containing sodium iodide (131-I) in parenteral dose form (medicinal product form)) |
| | • Silver sulfadiazine (e.g. 771756000 |Product containing silver sulfadiazine in cutaneous dose form (medicinal product form))] |
| Attribute: | |
| Has manufactured dose form | • Range: 736542009|Pharmaceutical dose form (dose form) - descendants that are groupers representing intended site only (e.g. 385268001 |Oral dose form (dose form)], 385287007 |Parenteral dose form (dose form))] |
| | • Cardinality: 1..1 |
| | • Exceptions: 385217004 |Conventional release gas for inhalation (dose form) may be used as manufactured dose form for Medicinal product form concepts. |
Naming Guidelines

**FSN**

Use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concepts that are selected as the attribute value, excluding the semantic tag. For multiple ingredient drug products, the active ingredients must be in alphabetical order and separated by the word “and”.

- Product containing `<Active ingredient FSN>` in `<Manufactured dose form FSN>` (medicinal product form)
- Product containing `<Active ingredient FSN>` and `<Active ingredient FSN>` in `<Manufactured dose form FSN>` (medicinal product form)
- Product containing `<Active ingredient FSN>` and `<Active ingredient FSN>` and `<Active ingredient FSN>` in `<Manufactured dose form FSN>` (medicinal product form)

Creation of MPF-containing concepts for all possible combinations of active ingredients contained in multiple ingredient products is not recommended at this time (no specific use case has been identified). For example, a product containing three active ingredients would only require creation of one MPF-containing concept. If any of the active ingredients is available as a single ingredient product, or as part of another multiple ingredient concept, then appropriate concepts would be created for those products.

Example:

- Product containing axitinib in oral dose form (medicinal product form)
- Product containing abacavir and lamivudine in oral dose form (medicinal product form)
- Product containing abacavir and lamivudine and zidovudine in oral dose form (medicinal product form)

**Preferred Term**

Use the following pattern for the PT; align naming and case sensitivity with the PT for the concept that is selected as the attribute value. For multiple ingredient drug products, the active ingredients must be in alphabetical order and separated by the word “and”.

- `<Active ingredient PT>`-containing product in `<Manufactured dose form PT>`
- `<Active ingredient PT>`- and `<Active ingredient PT>`-containing product in `<Manufactured dose form PT>`
- `<Active ingredient PT>`- and `<Active ingredient PT>`- and `<Active ingredient PT>`-containing product in `<Manufactured dose form PT>`

Example:

- Axitinib-containing product in oral dose form
- Abacavir- and lamivudine-containing product in oral dose form
- Abacavir- and lamivudine- and zidovudine-containing product in oral dose form

**Synonym**

Synonyms matching the FSN are not required.

---

**Exemplars**

The following illustrates the stated view for 773391004 |Product containing axitinib in oral dose form (medicinal product form)|
The following illustrates the inferred view for 773391004 |Product containing axitinib in oral dose form (medicinal product form)|:

The following illustrates the stated view for 767783007 |Product containing codeine and paracetamol in oral dose form (medicinal product form)|:
The following illustrates the inferred view for 767783007 |Product containing codeine and paracetamol in oral dose form (medicinal product form)|:

Medicinal Product Form containing only Concept

Overview

The Medicinal Product Form "containing only" (MPF-only) concept is an abstract representation of the active ingredient(s) and dose form intended site for a medicinal product. It means that the medicinal product must contain only the active ingredient(s) specified in the FSN but may also contain a modification of the active ingredient(s) specified in the FSN.
Example:
- Product containing only axitinib in oral dose form (medicinal product form)
- Product containing only abacavir and lamivudine in oral dose form (medicinal product form)

Modeling (stated view)
MPF-only concepts shall be modeled using the proximal primitive modeling pattern.

| Stated parent concept | 763158003|Medicinal product (product) |
|-----------------------|-----------------------------------------------|
| Semantic tag          | (medicinal product form) |
| Definition status     | 900000000000073002 |Sufficiently defined by necessary conditions definition status (core metadata concept) |
| Attribute:            | |
| Has active ingredient | |
| Range:                | 105590001|Substance (substance) - descendants only, excluding concepts representing structural groupers, dispositions, or combined substances |
| Cardinality:          | 1..∞ - there is no technical limit on the number of Has active ingredient attributes that may be added to a concept; a practical limit may be imposed at a later date. |
| For content in the International Release, this attribute value should represent the base ingredient, not a modification, unless explicitly identified as an exception. |
| Exceptions:           | |
| Chemical element with multiple modification (e.g. 422232005 |Calcium lactate gluconate (substance)) |
| Chloral hydrate (e.g. 778711000 |Product containing only chloral hydrate in oral dose form (medicinal product form)) |
| Liposome or lipid complex substances(e.g. 426490000 |Vincristine liposome (substance), 425953004 |Amphotericin B lipid complex (substance), 768664009 |Amphotericin B phospholipid complex (substance), 427544000 |Amphotericin B cholesteryl sulfate complex (substance)) |
| Exceptions:           | |
| Silver sulfadiazine (e.g. 864009007 |Product containing only silver sulfadiazine in cutaneous dose form (medicinal product form)) |

| Attribute:            | |
| Has manufactured dose form | |
| Range:                | 736542009|Pharmaceutical dose form (dose form) - descendants that are groupers representing intended site only (e.g. 385268001 |Oral dose form (dose form), 385287007 |Parenteral dose form (dose form)) |
| Cardinality:          | 1..1 |
| Exceptions:           | 385217004 |Conventional release gas for inhalation (dose form) may be used as manufactured dose form for Medicinal product form concepts.
### Attribute:

**Count of base of active ingredient**
- Concrete Type: Integer
- Range: >0..
- Cardinality: 1..1

### Naming Guidelines

#### FSN

Use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concepts that are selected as the attribute value, excluding the semantic tag. For multiple ingredient drug products, the active ingredients must be in alphabetical order and separated by the word “and”.

- Product containing only `<Active ingredient FSN>` in `<Manufactured dose form FSN>` (medicinal product form)
- Product containing only `<Active ingredient FSN>` and `<Active ingredient>` in `<Manufactured dose form FSN>` (medicinal product form)
- Product containing only `<Active ingredient FSN>` and `<Active ingredient FSN>` and `<Active ingredient FSN>` in `<Manufactured dose form FSN>` (medicinal product form)

Creation of MPF-only concepts for all possible combinations of active ingredients contained in multiple ingredient products is not recommended at this time (no specific use case has been identified). For example, a product containing three active ingredients would only require creation of one MPF-only concept. If any of the active ingredients is available as a single ingredient product, or as part of another multiple ingredient concept, then appropriate concepts would be created for those products.

**Example:**
- Product containing only axitinib in oral dose form (medicinal product form)
- Product containing only abacavir and lamivudine in oral dose form (medicinal product form)
- Product containing only abacavir and lamivudine and zidovudine in oral dose form (medicinal product form)

#### Preferred Term

Use the following pattern for the PT; align naming and case sensitivity with the PT for the concept that is selected as the attribute value. For multiple ingredient drug products, the active ingredients must be in alphabetical order and separated by the word “and”.

- `<Active ingredient PT>` only product in `<Manufactured dose form PT>`
- `<Active ingredient PT>`- and `<Active ingredient PT>`- only product in `<Manufactured dose form PT>`
- `<Active ingredient PT>`- and `<Active ingredient PT>`- and `<Active ingredient PT>` only product in `<Manufactured dose form PT>`

**Example:**
- Axitinib only product in oral dose form
- Abacavir and lamivudine only product in oral dose form
- Abacavir and lamivudine and zidovudine only product in oral dose form

#### Synonym

Synonyms matching the FSN are not required.

### Exemplars

The following illustrates the **stated** view for 773392006 |Product containing only axitinib in oral dose form (medicinal product form)|:
The following illustrates the **inferred** view for 773392006 |Product containing only axitinib in oral dose form (medicinal product form)|:
The following illustrates the inferred view for 772249008 |Product containing only abacavir and lamivudine in oral dose form (medicinal product form)|:
Clinical Drug containing precisely Concept - Discrete Dose Form

Overview

The Clinical Drug "containing precisely" (CD-precise) concept is an abstract representation of the precise active ingredient, basis of strength substance (BoSS), strength, and manufactured dose form of a drug product. It implies that the drug product must contain only the precise active ingredient(s) specified in the FSN.

Modeling for concepts representing discrete dose forms

CD-precise concepts shall be modeled using the proximal primitive modeling pattern.

CD-precise concepts representing discrete dose form (e.g. tablets, capsules, pessaries, suppositories, sachets, ampules or vials containing solid dose forms such as powders or granules, and metered dose delivery products such as inhalers and spray) will be modeled using presentation strength attributes; concentration strength attributes are not allowed for these concepts in the International Release.

Example:

- Product containing precisely abacavir 300 milligram/1 each conventional release oral tablet (clinical drug)
- Product containing precisely abacavir 600 milligram and lamivudine 300 milligram/1 each conventional release oral tablet (clinical drug)
- Product containing precisely afatinib (as afatinib dimaleate) 30 milligram/1 each conventional release oral tablet (clinical drug)
- Product containing precisely aztreonam 500 milligram/1 vial powder for conventional release solution for injection (clinical drug)
- Product containing precisely flucloxacillin (as flucloxacillin sodium) 250 milligram/1 vial powder for conventional release solution for injection (clinical drug)
- Product containing precisely budesonide 200 microgram/actuation conventional release powder for inhalation (clinical drug)

The following exceptions have been identified.

- Concepts with product strength that is "not equal to"
  - These concepts will be modeled as primitive, with attributes added as described in the following sections, except that strength numerator attributes will not be added.
  - These concepts will have the appropriate Medicinal Product Form-only (MPF-only) concept as an inferred parent.

| Stated parent concept | 763158003|Medicinal product (product) |
|-----------------------|------------------|
| Semantic tag          | (clinical drug)   |
| Definition status     | 900000000000073002 |Sufficiently defined by necessary conditions definition status (core metadata concept)|
|                        |                      |
| Exceptions:           |                      |
| - Concepts with product strength that is "not equal to" (e.g. with product strength expressed as a range, greater than, or less than) will have definition status 900000000000074008|Necessary but not sufficient concept definition status (core metadata concept)|
### Attribute: Has manufactured dose form

- **Range:** 736542009 | Pharmaceutical dose form (dose form) - descendants only  
  - **NOTE:** While the allowed range for this attribute is broader, the CD-precise discrete dose form concepts should only use descendants of 736542009 | Pharmaceutical dose form (dose form), excluding grouper concepts based on intended site (e.g. 740596000 | Cutaneous dose form (dose form)), 385268001 | Oral dose form (dose form)).
  - **Cardinality:** 1..1

The attribute value should represent the **manufactured** dose form

**Notes:**

- There are known inconsistencies with respect to the use of "injection" or "infusion"; the need for content in the International Release to be consistent and predictable is recognized. Further refinements will be made in a future release based on feedback from active users of the content and concept model at a future date.
- Concepts with dose form of drops will be modeled in the International Release with Pharmaceutical dose form concepts that do not specify the basic dose form (e.g. 385018001 | Conventional release oral drops (dose form)).
  - More specific Pharmaceutical dose form concepts (e.g. 385019009 | Conventional release solution for oral drops (dose form)) will be retained in the International Release but will not be used for modeling Clinical drug concepts; they can be used for modeling in national extensions.
- Powder for oral suspension, solution, etc. may be modeled using concentration strength and the administrable dose form (e.g. 1145409004 | Product containing precisely amoxicillin 25 milligram/1 milliliter and clavulanic acid (as clavulanate potassium) 6.25 milligram/1 milliliter conventional release oral suspension (clinical drug)).
- Clinical drug concepts in the International Release should not be modeled using dose forms for "lyophilized powder" because it is not a clinically relevant distinction at the Clinical drug level.
  - Pharmaceutical dose forms referring to "lyophilized powder" will be retained in the International Release but will not be used for modeling Clinical drug concepts; they can be used for modeling in national extensions.

### Attribute: Has unit of presentation

- **Range:** 732935002 | Unit of presentation (unit of presentation) - descendants only
  - **Cardinality:** 1..1

### Attribute: Count of base of active ingredient (attribute)

- **Concrete Type:** Integer
- **Range:** >=0.. 
- **Cardinality:** 1..1

### Relationship group

One relationship group containing one instance of each of the following attributes is required for each precise active ingredient.

- **Has precise active ingredient**
  - **Range:** 105590001 | Substance (substance) - descendants only, excluding concepts representing structural groupers, dispositions, or combined substances
  - **Cardinality:** 1..1 per relationship group

**Notes:**

- The PAI cannot be modeled as a substance hydrate or solvate unless the BoSS is expressed as a hydrate or solvate.
<table>
<thead>
<tr>
<th>Attribute</th>
<th>Value</th>
</tr>
</thead>
</table>
| Has basis of strength sub stance| - Range: 105590001|Substance (substance) - descendants only, excluding concepts representing structural groupers, dispositions, or combined substances  
- Cardinality: 1..1 per relationship group |
| Has presentation strength numerator value | - Concrete Type: Decimal  
- Range: >#0..  
- Cardinality: 1..1 per relationship group  
Expression of product strength in metric units is preferred. To avoid semantically equivalent concepts, product strength for metric units are normalized as follows:  
- Use milligram if value is <1000; if ≥ then convert to gram  
- Use microgram if value is <1000; if ≥ then convert to milligram  
- Use nanogram if value is <1000; if ≥ then convert to microgram  
- Use picogram if value is <1000; if ≥ then convert to nanogram  
To avoid semantically equivalent concepts, product strength for units are normalized as follows:  
- Use million unit if value is >1000000 unit  
The following units are not allowed unless specifically noted as an exception:  
- 408165007 |Mega u (qualifier value)|  
Repeating decimals are rounded to three decimal places (with 5 and above rounded up and 4 and below rounded down); for example:  
- 769539009 |833.333 (qualifier value)|  
- 769538001 |666.667 (qualifier value)|  
- 769339008 |104.167 (qualifier value)| |
| Has presentation strength denominator unit | - Range: 767524001 |Unit of measure (qualifier value)| - descendants only  
- Cardinality: 1..1 per relationship group |
| Has presentation strength denominator value | - Concrete Type: Decimal  
- Range: >#0..  
- Cardinality: 1..1 per relationship group  
- For this pattern the attribute value must be 38112003 |1 (qualifier value)|  
Note: the denominator strength value is required for concepts in the International Release even if the value = 1 because including denominators for only some concepts negatively affects the classification results. |
| Has presentation strength denominator unit | - Range: 767524001 |Unit of measure (qualifier value)| - descendants only  
- NOTE: While the allowed range for this attribute is broader, the CD-precise concepts representing discrete dose forms should only use descendants of 732935002|Unit of presentation (unit of presentation).  
- Cardinality: 1..1 per relationship group |
### Naming

Use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concepts that are selected as the attribute values, excluding the semantic tag. For multiple ingredient drug products, the BoSS must be in alphabetical order and separated by the word "and".

Where **Precise active ingredient** = BoSS and **Unit of presentation** = discrete solid dose form (e.g. capsule, lozenge, pessary, suppository, tablet):

- Product containing precisely `<BoSS FSN>` `<Presentation strength numerator value FSN>` `<Presentation strength numerator unit FSN>` / `<Presentation strength denominator value FSN>` each `<Manufactured dose form FSN>` (clinical drug)

Example:

- Product containing precisely abacavir 300 milligram/1 each conventional release oral tablet (clinical drug)
- Product containing precisely abacavir 600 milligram and lamivudine 300 milligram/1 each conventional release oral tablet (clinical drug)
- Product containing precisely atropine sulfate 600 microgram/1 each conventional release oral tablet (clinical drug)
- Product containing precisely codeine sulfate 15 milligram/1 each conventional release oral tablet (clinical drug)

Where **Precise active ingredient** <> BoSS and **Unit of presentation** = discrete solid dose form (e.g. capsule, lozenge, pessary, suppository, tablet):

- Product containing precisely `<BoSS FSN>` (as `<Precise active ingredient FSN>`) `<Presentation strength numerator value FSN>` `<Presentation strength numerator unit FSN>` / `<Presentation strength denominator value FSN>` each `<Manufactured dose form FSN>` (clinical drug)

Example:

- Product containing precisely doxazosin (as doxazosin mesilate) 4 milligram/1 each conventional release oral tablet (clinical drug)
- Product containing precisely disopyramide (as disopyramide phosphate) 150 milligram/1 each prolonged-release oral tablet (clinical drug)

Where **Precise active ingredient** = BoSS and **Unit of presentation** = other discrete dose form (e.g. actuation, vial, sachet)

- Product containing precisely `<BoSS FSN>` `<Presentation strength numerator value FSN>` `<Presentation strength numerator unit FSN>` / `<Presentation strength denominator value FSN>` `<Presentation strength denominator unit FSN>` `<Manufactured dose form FSN>` (clinical drug)

Example:

- Product containing precisely aztreonam 500 milligram/1 vial powder for conventional release solution for injection (clinical drug)
- Product containing precisely budesonide 200 microgram/1 actuation conventional release powder for inhalation (clinical drug)

Where **Precise active ingredient** <> BoSS and **Unit of presentation** = other discrete dose form (e.g. actuation, vial, sachet)
• Product containing precisely <BoSS FSN> (as <Precise active ingredient FSN>) <Presentation strength numerator value FSN> <Presentation strength numerator unit FSN>/<Presentation strength denominator value FSN> <Presentation strength denominator unit FSN> Manufactured dose form FSN> (clinical drug)

Example

• Product containing precisely flucloxacillin (as flucloxacillin sodium) 250 milligram/1 vial powder for conventional release solution for injection (clinical drug)
• Product containing precisely buserelin (as buserelin acetate) 100 microgram/1 actuation conventional release nasal spray (clinical drug)
Use the following pattern for the PT; align naming and case sensitivity with the PT for the concepts that are selected as the attribute values. For multiple ingredient drug products, the BoSS must be in alphabetical order and separated by the word “and”.

Where BoSS = Precise active ingredient:

- <BoSS PT> <Presentation strength numerator value PT> <Presentation strength numerator unit PT> <Manufactured dose form PT>

Example:

- Abacavir 300 mg oral tablet
- Abacavir 600 mg and lamivudine 300 mg oral tablet
- Atropine sulfate 600 microgram oral tablet
- Codeine sulfate 15 mg oral tablet
- Aztreonam 500 mg powder for solution for injection vial
- Budesonide 200 microgram/actuation powder for inhalation
- Buserelin (as buserelin acetate) 100 microgram/actuation nasal spray

Where BoSS <> Precise active ingredient:

- <BoSS PT> (as <Precise active ingredient PT>) <Presentation strength numerator value PT> <Presentation strength numerator unit PT> <Manufactured dose form PT>

Example:

- US PT: Doxazosin (as doxazosin mesylate) 4 mg oral tablet
- GB PT: Doxazosin (as doxazosin mesylate) 4 mg oral tablet
- US/GB PT: Disopyramide (as disopyramide phosphate) 150 mg prolonged-release oral tablet
- US PT: Floxacillin (as floxacillin sodium) 250 mg powder for solution for injection vial
- GB PT: Flucloxacillin (as flucloxacillin sodium) 250 mg powder for solution for injection vial

Exception:

- The following units of measure should not be abbreviated in descriptions; they should always be spelled out:
  - international unit
  - microequivalent
  - microgram
  - microliter (with GB spelling microlitre)
  - million unit
  - nanogram
  - picogram
  - unit

Synonyms

Synonyms matching the FSN are not required.

Exemplars

The following illustrates the stated view for 318783003 |Product containing precisely doxazosin (as doxazosin mesilate) 4 milligram/1 each conventional release oral tablet (clinical drug)|:
The following illustrates the inferred view for 318783003 |Product containing precisely doxazosin (as doxazosin mesilate) 4 milligram/1 each conventional release oral tablet (clinical drug)|:
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(2021-10-02)

The following illustrates the stated view for 783301004 |Product containing precisely abacavir (as abacavir sulfate) 600 milligram and lamivudine 300 milligram/1 each conventional release oral tablet (clinical drug)|:
The following illustrates the **inferred** view for 783301004 |Product containing precisely abacavir (as abacavir sulfate) 600 milligram and lamivudine 300 milligram/1 each conventional release oral tablet (clinical drug)|:
Clinical Drug containing precisely Concept - Continuous Dose Form

Overview
The Clinical Drug "containing precisely" (CD-precise) concept is an abstract representation of the precise active ingredient, basis of strength substance (BoSS), strength, and manufactured dose form of a drug product. It implies that the drug product must contain only the precise active ingredient(s) specified in the FSN.

Modeling for concepts representing continuous dose forms
CD-precise concepts shall be modeled using the proximal primitive modeling pattern.
CD-precise concepts representing continuous dose form (e.g. solutions, suspensions, creams, ointments, patches) will be modeled using concentration strength attributes; presentation strength attributes are not allowed for these concepts in the International Release.

Example:
- Product containing precisely zidovudine 10 milligram/1 milliliter conventional release oral solution (clinical drug)
- Product containing precisely amikacin (as amikacin sulfate) 250 milligram/1 milliliter conventional release solution for injection (clinical drug)
- Product containing precisely clotrimazole 10 milligram/1 gram conventional release cutaneous cream (clinical drug)
- Product containing precisely mupirocin (as mupirocin calcium) 20 milligram/1 gram conventional release nasal ointment (clinical drug)
- Product containing precisely buprenorphine 70 microgram/1 hour prolonged-release transdermal patch (clinical drug)

The following exceptions have been identified.
- Concepts with product strength that is "not equal to"
  - These concepts will be modeled as primitive, with attributes added as described in the following sections, except that strength numerator attributes will not be added.
  - These concepts will have the appropriate Medicinal Product Form-only (MPF-only) concept as an inferred parent.

| Stated parent concept | 763158003|Medicinal product (product) |
| Semantic tag | (clinical drug) |
| Definition status | 900000000000073002 |Sufficiently defined by necessary conditions definition status (core metadata concept)|

Exceptions:
- Concepts with product strength that is "not equal to" (e.g. with product strength expressed as a range, greater than, or less than) will have definition status 900000000000074008 | Necessary but not sufficient concept definition status (core metadata concept) |
<table>
<thead>
<tr>
<th>Attribute:</th>
<th>Has manufactured dose form</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Range:</strong></td>
<td>736542009</td>
</tr>
<tr>
<td>• NOTE: While the allowed range for this attribute is broader, the CD-precise continuous dose from concepts should only use descendants of 736542009</td>
<td>Pharmaceutical dose form (dose form), excluding grouper concepts based on intended site (e.g. 740596000</td>
</tr>
<tr>
<td>• <strong>Cardinality:</strong></td>
<td>1..1</td>
</tr>
<tr>
<td>The attribute value should represent the <strong>manufactured</strong> dose form</td>
<td></td>
</tr>
</tbody>
</table>

**Notes:**

- There are known inconsistencies with respect to the use of "injection" or "infusion"; the need for content in the International Release to be consistent and predictable is recognized. Further refinements will be made in a future release based on feedback from active users of the content and concept model at a future date.  
- Concepts with dose form of drops will be modeled in the International Release with general Pharmaceutical dose form concepts (e.g. 385018001 | Conventional release oral drops (dose form)|).  
  - More specific Pharmaceutical dose form concepts (e.g. 385019009 | Conventional release solution for oral drops (dose form)|) will be available for use to model content in national extensions.  
  - Powder for oral suspension, solution, etc. may be modeled using concentration strength and the administrable dose form (e.g. 1145409004 | Product containing precisely amoxicillin 25 milligram/1 milliliter and clavulanic acid (as clavulanate potassium) 6.25 milligram/1 milliliter conventional release oral suspension (clinical drug)|)

<table>
<thead>
<tr>
<th>Attribute:</th>
<th>Count of base of active ingredient (attribute)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Concrete Type:</strong></td>
<td>Integer</td>
</tr>
<tr>
<td><strong>Range:</strong></td>
<td>&gt;0..</td>
</tr>
<tr>
<td><strong>Cardinality:</strong></td>
<td>1..1</td>
</tr>
</tbody>
</table>

**Relationship group**  
One relationship group containing one instance of each of the following attributes is required for each precise active ingredient.

- **Has precise active ingredient**  
  - **Range:** 105590001 | Substance (substance) - descendants only, excluding concepts representing structural groupers, dispositions, or combined substances |
  - **Cardinality:** 1..1 per relationship group  
  - Notes:  
    - The PAI cannot be modeled as a substance hydrate or solvate unless the BoSS is expressed as a hydrate or solvate.

- **Has basis of strength substance**  
  - **Range:** 105590001 | Substance (substance) - descendants only, excluding concepts representing structural groupers, dispositions, or combined substances |
  - **Cardinality:** 1..1 per relationship group
• Has concentration strength numerator value
  • Concrete Type: Decimal
  • Range: >0..
  • Cardinality: 1..1 per relationship group
  Expression of product strength in metric units is preferred.
  To avoid semantically equivalent concepts, product strength for metric units are normalized as follows:
  • Use milligram if value is <1000; if ≥ then convert to gram
  • Use microgram if value is <1000; if ≥ then convert to milligram
  • Use nanogram if value is <1000; if ≥ then convert to microgram
  • Use picogram if value is <1000; if ≥ then convert to nanogram
  To avoid semantically equivalent concepts, product strength for units are normalized as follows:
  • Use million unit if value is ≥1000000 unit

The following units are not allowed unless specifically noted as an exception:

  • 408165007 [Mega u (qualifier value)]

Repeating decimals are rounded to three decimal places (with 5 and above rounded up and 4 and below rounded down); for example:

  • 769539009 [833.333 (qualifier value)]
  • 769538001 [666.667 (qualifier value)]
  • 769339008 [104.167 (qualifier value)]

• Has concentration strength numerator unit
  • Range: 767524001 [Unit of measure (qualifier value)] - descendants only
  • Cardinality: 1..1 per relationship group

• Has concentration strength denominator value
  • Concrete Type: Decimal
  • Range: >0..
  • Cardinality: 1..1 per relationship group
  • For this pattern the attribute value must be 38112003 [1 (qualifier value)]
  • Exceptions: none identified

Note: the denominator strength value is required for concepts in the International Release even though the value = 1 because including denominators for only some concepts negatively affects the classification results.

• Has concentration strength denominator unit
  • Range: 767524001 [Unit of measure (qualifier value)] - descendants only
  • Cardinality: 1..1 per relationship group

  • For Clinical drug concepts with liquid dose form, the denominator unit should be 258773002 [Milliliter (qualifier value)].
  • For Clinical drug concepts with semi-solid dose form, the denominator unit should be 258682000 [gram (qualifier value)] for weight/weight concentration and 258773002 [Milliliter (qualifier value)] for weight/volume concentration.
## Naming

<table>
<thead>
<tr>
<th>FSN</th>
<th>Use the following pattern for the FSN; align naming and case sensitivity with the FSN for the concepts that are selected as the attribute values, excluding the semantic tag. For multiple ingredient drug products, the BoSS must be in alphabetical order and separated by the word &quot;and&quot;.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td><strong>Where Precise active ingredient = BoSS:</strong></td>
</tr>
<tr>
<td></td>
<td>• Product containing precisely <code>&lt;BoSS FSN&gt;</code> <code>&lt;Concentration strength numerator value FSN&gt;</code> <code>&lt;Concentration strength numerator unit FSN&gt;</code>/ <code>&lt;Concentration strength denominator value FSN&gt;</code> <code>&lt;Concentration strength denominator unit FSN&gt;</code> <code>&lt;Manufactured dose form FSN&gt;</code> (clinical drug)</td>
</tr>
<tr>
<td></td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td>• Product containing precisely amitriptyline hydrochloride 5 milligram/1 milliliter conventional release oral solution (clinical drug)</td>
</tr>
<tr>
<td></td>
<td>• Product containing precisely buprenorphine 70 microgram/1 hour prolonged-release transdermal patch (clinical drug)</td>
</tr>
<tr>
<td></td>
<td>• Product containing precisely clotrimazole 10 milligram/1 gram conventional release cutaneous cream (clinical drug)</td>
</tr>
<tr>
<td></td>
<td>• Product containing precisely zidovudine 10 milligram/1 milliliter conventional release oral solution (clinical drug)</td>
</tr>
<tr>
<td></td>
<td>• Product containing precisely lopinavir 80 milligram/1 milliliter and ritonavir 20 milligram/1 milliliter conventional release oral solution (clinical drug)</td>
</tr>
<tr>
<td></td>
<td><strong>Where Precise active ingredient &lt;&gt; BoSS:</strong></td>
</tr>
<tr>
<td></td>
<td>• Product containing precisely <code>&lt;BoSS FSN&gt;</code> (as <code>&lt;Precise active ingredient FSN&gt;</code>) <code>&lt;Concentration strength numerator value FSN&gt;</code> <code>&lt;Concentration strength numerator unit FSN&gt;</code>/ <code>&lt;Concentration strength denominator value FSN&gt;</code> <code>&lt;Concentration strength denominator unit FSN&gt;</code> <code>&lt;Manufactured dose form FSN&gt;</code> (clinical drug)</td>
</tr>
<tr>
<td></td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td>• Product containing precisely amikacin (as amikacin sulfate) 250 milligram/1 milliliter conventional release solution for injection (clinical drug)</td>
</tr>
<tr>
<td></td>
<td>• Product containing precisely mupirocin (as mupirocin calcium) 20 milligram/1 gram conventional release nasal ointment (clinical drug)</td>
</tr>
</tbody>
</table>
| Preferred Term | Use the following pattern for the PT; align naming and case sensitivity with the PT for the concepts that are selected as the attribute values. For multiple ingredient drug products, the BoSS must be in alphabetical order and separated by the word “and”.

Where Single ingredient with BoSS = Precise active ingredient:

- `<BoSS PT> <Concentration strength numerator value PT> <Concentration strength numerator unit PT> / <Concentration strength denominator value PT> <Concentration strength denominator unit PT> <Manufactured dose form PT>`

Example:

- Amitriptyline hydrochloride 5 mg/mL oral solution
- Buprenorphine 70 microgram/hour prolonged-release transdermal patch
- Clotrimazole 10 mg/g cutaneous cream
- Zidovudine 10 mg/mL oral solution
- Lopinavir 80 mg/mL and ritonavir 20 mg/mL oral solution

Where Single ingredient with BoSS <> Precise active ingredient:

- `<BoSS PT> (as <Precise active ingredient PT>) <Concentration strength numerator value PT> <Concentration strength numerator unit PT> / <Concentration strength denominator value PT> <Concentration strength denominator unit PT> <Manufactured dose form PT>`

Example:

- Amikacin (as amikacin sulfate) 250 mg/mL solution for injection
- Mupirocin (as mupirocin calcium) 20 mg/g nasal ointment

Exception:

- The following units of measure should not be abbreviated in descriptions; they should always be spelled out:
  - international unit
  - microequivalent
  - microliter
  - microgram
  - microliter (with GB spelling microlitre)
  - million unit
  - nanogram
  - picogram
  - unit

| Synonyms | Synonyms matching the FSN are not required. Synonyms converting metric units to percent or parts per millions may be included for medical gas concepts (e.g. Helium 79% and oxygen 21% gas for inhalation, Helium 790,000 ppm and oxygen 210,000 ppm gas for inhalation) |

Exemplars

The following illustrates the stated view for 781834001 |Product containing precisely digoxin 250 microgram/1 milliliter conventional release solution for injection (clinical drug)|:
The following illustrates the *inferred* view for 781834001 [Product containing precisely digoxin 250 microgram/1 milliliter conventional release solution for injection (clinical drug)]:
The following illustrates the stated view for 769821007 |Product containing precisely digoxin 50 microgram/1 milliliter conventional release oral solution (clinical drug)|:
The following illustrates the *inferred* view for 769821007 [Product containing precisely digoxin 50 microgram/1 milliliter conventional release oral solution (clinical drug)]:

![Diagram of inferred view for 769821007](image-url)
The following illustrates the stated view for 396279000 |Product containing precisely betamethasone (as betamethasone valerate) 1 milligram/1 gram conventional release cutaneous ointment (clinical drug)|:
The following illustrates the inferred view for 396279000 |Product containing precisely betamethasone (as betamethasone valerate) 1 milligram/1 gram conventional release cutaneous ointment (clinical drug)|:
The following illustrates the stated view for 769514000 |Product containing precisely buprenorphine 70 microgram/1 hour prolonged-release transdermal patch (clinical drug)|:
The following illustrates the inferred view for 769514000 |Product containing precisely buprenorphine 70 microgram/1 hour prolonged-release transdermal patch (clinical drug)|:
Vaccine Products in the Medicinal Product Hierarchy

The following sections apply to the vaccine product concepts in the |Medicinal product| hierarchy in the International Release.

In the International Release, vaccine products are those concepts with attribute Plays role = Active immunity stimulant therapeutic role (role). Products that provide passive immunity should be modeled using the general Medicinal product guidelines.

Note: Modeling "antigen" as a role or disposition may be considered in the future but is out of scope at this time.

Vaccine Product Top Level Groupers

Overview

The following high level vaccine-related grouper concepts will be included in the |Medicinal product| hierarchy.

- 787859002 |Vaccine product (medicinal product) |
- 836368004 |Vaccine product containing bacteria antigen (medicinal product) |
- 836369007 |Vaccine product containing virus antigen (medicinal product) |
- 863950005 |Vaccine product containing bacteria and virus antigens (medicinal product) |
### Modeling (stated view)

<table>
<thead>
<tr>
<th><strong>Stated parent concept</strong></th>
<th>763158003</th>
<th>Medicinal product (product)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Semantic tag</strong></td>
<td>(medicinal product)</td>
<td></td>
</tr>
<tr>
<td><strong>Definition status</strong></td>
<td>900000000000073002</td>
<td>Sufficiently defined by necessary conditions definition status (core metadata concept)</td>
</tr>
<tr>
<td><strong>Attribute:</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Has active ingredient</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range:</td>
<td>&lt;=105590001</td>
<td>Substance (substance)</td>
</tr>
<tr>
<td>Cardinality:</td>
<td>0..*</td>
<td></td>
</tr>
<tr>
<td><strong>Attribute:</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td><strong>Plays role</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Range:</td>
<td>&lt;=766940004</td>
<td>Role (role)</td>
</tr>
<tr>
<td>Cardinality:</td>
<td>0..*</td>
<td></td>
</tr>
<tr>
<td><strong>NOTE:</strong> While the allowed range is broader, top level vaccine-related grouper concepts should have one and only one</td>
<td>Plays role attribute with attribute value = 31833100221102</td>
<td>Active immunity stimulant therapeutic role (role).</td>
</tr>
</tbody>
</table>

### Naming Guidelines

| **FSN**                   | Use the following pattern for the FSN; align naming and case sensitivity with the PT for the concept that is selected as the attribute value for the 127489000 | Has active ingredient (attribute). |
|---------------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
|                           | • Vaccine product containing `<Active ingredient PT excluding "antigen"> antigen (medicinal product)`                                                                                          |
|                           | • Vaccine product containing `<Active ingredient PT excluding "antigen"> and <Active ingredient PT excluding "antigen"> antigens (medicinal product)`                                    |
| **Exception:**            |                                                                                                                                                                                            |
|                           | • Top level grouper 787859002 | Vaccine product (medicinal product) | does not have a Has active ingredient attribute. |
| **Example:**              |                                                                                                                                                                                            |
|                           | • Vaccine product (medicinal product)                                                                                                                                                          |
|                           | • Vaccine product containing bacteria antigen (medicinal product)                                                                                                                           |
|                           | • Vaccine product containing virus antigen (medicinal product)                                                                                                                            |
|                           | • Vaccine product containing bacteria and virus antigens (medicinal product)                                                                                                               |
### Preferred Term

Use the following pattern for the PT; align naming and case significance with the PT for the concept that is selected as the attribute value for the 127489000 |Has active ingredient (attribute)|.

- `<Active ingredient PT excluding "antigen">-containing vaccine product`
- `<Active ingredient PT excluding "antigen">- and `<Active ingredient PT excluding "antigen"> antigens-containing vaccine product`

**Exception:**

- Top level grouper 787859002 |Vaccine product (medicinal product)| does not have a Has active ingredient attribute.

**Example:**

- Vaccine product
- Bacteria antigen-containing vaccine product
- Virus antigen-containing vaccine product
- Bacteria- and virus antigens-containing vaccine product

### Synonyms

Synonyms matching the FSN are not required.

### Exemplars

The following illustrates the **stated** view for 787859002 |Vaccine product (product)|:

- **787859002 Vaccine product (medicinal product)**
  - **765188003 Medicinal product (product)**
    - **766090001 Plays role (attribute)**
      - **318331000221102 Active immunity stimulant therapeutic role (role)**

The following illustrates the **inferred** view for 787859002 |Vaccine product (product)|:

- **787859002 Vaccine product (medicinal product)**
  - **763067004 Medicinal product categorized by therapeutic role (product)**
    - **766090001 Plays role (attribute)**
      - **318331000221102 Active immunity stimulant therapeutic role (role)**

The following illustrates the **stated** view for 836368004 |Vaccine product containing bacteria antigen (medicinal product)|:
The following illustrates the inferred view for 836368004 |Vaccine product containing bacteria antigen (medicinal product)|:

![Diagram 1]

The following illustrates the stated view for 863950005 |Vaccine product containing bacteria and virus antigens (medicinal product)|:

![Diagram 2]

The following illustrates the inferred view for 863950005 |Vaccine product containing bacteria and virus antigens (medicinal product)|:
Vaccine Product containing Concepts

Overview

The Vaccine Product "containing" concept is an abstract representation of the active ingredient(s) in a vaccine product. It means that the vaccine product must contain the active ingredient(s) specified in the FSN but may also contain a modification of the active ingredient(s) specified in the FSN or may contain additional active ingredient(s).

Example:

- 836374004 |Vaccine product containing Hepatitis B virus antigen (medicinal product)|
- 836389008 |Vaccine product containing Vaccinia virus antigen (medicinal product)|

Both vaccine product "containing" and vaccine product "containing only" concepts may be created for products that only have one active ingredient (e.g. 836374004 |Vaccine product containing Hepatitis B virus antigen (medicinal product)| and 871822003 |Vaccine product containing only Hepatitis B virus antigen (medicinal product)|).

Vaccine product "containing" concepts are not created for multiple ingredient vaccine products; vaccine product "containing only" concepts are created for multiple ingredient vaccine products.

Modeling (stated view)

<table>
<thead>
<tr>
<th>Stated parent concept</th>
<th>763158003</th>
<th>Medicinal product (product)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Semantic tag</td>
<td>(medicinal product)</td>
<td></td>
</tr>
<tr>
<td>Definition status</td>
<td>90000000000073002</td>
<td>Sufficiently defined by necessary conditions definition status (core metadata concept)</td>
</tr>
</tbody>
</table>
Attribute: Has active ingredient
- Range: 105590001|Substance (substance) - descendants only, excluding concepts representing structural groupers, dispositions, or combined substances
- Cardinality: 1..*
  - While the allowed range is broader, Vaccine product "containing" concepts in the International Release should have one and only one [Has active ingredient] attribute.
  - For content in the International Release, this attribute value should represent the organism antigen, not a modification or subtype, unless explicitly identified as an exception.
  - Exceptions: Vaccine product containing concepts for the following substance subtypes are included (to support vaccination certificates):
    - 161000221102 |Antigen of Corynebacterium diphtheriae toxoid (substance)|
    - 551000221106 |Antigen of Clostridium tetani toxoid (substance)|

Attribute: Plays role
- Range: <=766940004 |Role (role)|
- Cardinality: 0..*
  - NOTE: While the allowed range is broader, Vaccine product "containing" concepts should have one and only one [Plays role] attribute with attribute value = 318331000221102 |Active immunity stimulant therapeutic role (role)|.

Naming Guidelines
FSN
Use the following pattern for the FSN; align naming and case sensitivity with the PT for the concept that is selected as the attribute value for the 127489000 |Has active ingredient (attribute)|. For multiple ingredient vaccine products, the active ingredients must be listed in alphabetical order, separated by the word "and", and the word "antigen" will be omitted. For concepts where all active ingredients are virus, the word "virus" may be omitted and added before "antigens".
- Vaccine product containing <Active ingredient PT> (medicinal product)
  Example:
  - Vaccine product containing Hepatitis B virus antigen (medicinal product)
  - Vaccine product containing Haemophilus influenzae type B antigen (medicinal product)

Preferred Term
Use the following pattern for the PT; align naming and case significance with the PT for the concept that is selected as the attribute value for the 127489000 |Has active ingredient (attribute)|. For multiple ingredient vaccine products, the active ingredients must be listed in alphabetical order, separated by the word "and", and the word "antigen" will be omitted. For concepts where all active ingredients are virus, the word "virus" may be omitted and added before "antigens".
- <Active ingredient PT>-containing vaccine product
  Example:
  - Hepatitis B virus antigen-containing vaccine product
  - Haemophilus influenzae type B antigen-containing product

Synonyms
Synonyms matching the FSN are not required

Exemplars
The following illustrates the stated view for 836374004 |Vaccine product containing Hepatitis B virus antigen (medicinal product)|:
The following illustrates the inferred view for 836374004 |Vaccine product containing Hepatitis B virus antigen (medicinal product)|:

Vaccine Product containing only Concepts

Overview

The Vaccine Product "only" concept is an abstract representation of the active ingredient(s) in a vaccine product. It means that the vaccine product must contain only the active ingredient(s) specified in the FSN but may also contain a modification of the active ingredient(s) specified in the FSN. The vaccine product "containing only" may be sufficient to serve as an interoperability layer or to support prescribing use cases.

Example:

- [Vaccine product containing only Hepatitis B virus antigen (medicinal product)]
- [Vaccine product containing only Vaccinia virus antigen (medicinal product)]
- [Vaccine product containing only Hepatitis A and Hepatitis B virus antigens (medicinal product)]
- [Vaccine product containing only Bordetella pertussis and Clostridium tetani and Corynebacterium diphtheriae antigens (medicinal product)]

Note: Both vaccine product "containing" and vaccine product "containing only" concepts may be created for products that only have one active ingredient (e.g. 836374004 |Vaccine product containing Hepatitis B virus antigen (medicinal product)| and 871822003 |Vaccine product containing only Hepatitis B virus antigen (medicinal product)|). Vaccine product "containing" concepts are not created for multiple ingredient vaccine products; vaccine product "containing only" concepts are created for multiple ingredient vaccine products.

Note: Modeling and terming for vaccines that have variable composition (e.g. influenza that may be specific to a year or hemisphere) will be addressed at a future date when use cases and requirements are better understood.
## Modeling (stated view)

<table>
<thead>
<tr>
<th>Stated parent concept</th>
<th>763158003</th>
<th>Medicinal product (product)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exceptions: none identified</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(medicinal product)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Definition status</td>
<td>90000000000073002</td>
</tr>
<tr>
<td>Exceptions: none identified</td>
<td></td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attribute: Has active ingredient</th>
</tr>
</thead>
<tbody>
<tr>
<td>Range: 105590001</td>
</tr>
<tr>
<td>Cardinality: 1..* - there is no technical limit on the number of Has active ingredient attributes that may be added to a concept; a practical limit may be imposed at a later date.</td>
</tr>
<tr>
<td>Exceptions: none identified</td>
</tr>
</tbody>
</table>

For content in the International Release, this attribute value should represent either the organism antigen, or the organism antigen(s), including modifications or subtypes, that are contained in a manufactured product.

<table>
<thead>
<tr>
<th>Attribute: Ingredient qualitative strength</th>
</tr>
</thead>
<tbody>
<tr>
<td>Range: &lt; 1149484003</td>
</tr>
<tr>
<td>Cardinality: 0..* - there is no technical limit on the number of Has ingredient characteristic attributes that may be added to a concept; a practical limit may be imposed at a later date.</td>
</tr>
<tr>
<td>Note: While the attribute range is large for this attribute, there are only as small number of concepts that are viable values for this attribute. The range may be refined or narrowed at a later date.</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attribute: Has target population</th>
</tr>
</thead>
<tbody>
<tr>
<td>Range: &lt; 27821000087106</td>
</tr>
<tr>
<td>Cardinality: 0..1</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attribute: Plays role</th>
</tr>
</thead>
<tbody>
<tr>
<td>Range: &lt;766940004</td>
</tr>
<tr>
<td>Exceptions: none identified</td>
</tr>
<tr>
<td>Cardinality: 0..*</td>
</tr>
<tr>
<td>NOTE: While the allowed range is broader, Vaccine product &quot;containing&quot; concepts should have one and only one</td>
</tr>
<tr>
<td>Exceptions: none identified</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Attribute: Count of base of active ingredient</th>
</tr>
</thead>
<tbody>
<tr>
<td>Range: 260299005</td>
</tr>
<tr>
<td>Cardinality: 1..1</td>
</tr>
</tbody>
</table>

For content in the International Release, this attribute value should represent the total number of discrete active ingredients, excluding modifications or subtypes.

<table>
<thead>
<tr>
<th>Attribute: Count of active ingredient</th>
</tr>
</thead>
<tbody>
<tr>
<td>Range: 260299005</td>
</tr>
<tr>
<td>Cardinality: 0..1</td>
</tr>
</tbody>
</table>

For content in the International Release, this attribute value should represent the total number of discrete active ingredients, including modifications or subtypes. This attribute is added only if the number of active ingredients is different from the number of base of active ingredients.
-called "antigens". For concepts where all active ingredients are virus, the word "virus" may be omitted and added before "antigens".

- Vaccine product containing only <Active ingredient PT> (medicinal product)
- Vaccine product containing only <Active ingredient PT> and <Active ingredient PT> antigens (medicinal product)
- Vaccine product containing only <Active ingredient PT> and <Active ingredient PT> and <Active ingredient PT> antigens (medicinal product)

Example:

- Vaccine product containing only Hepatitis B virus antigen (medicinal product)
- Vaccine product containing only Hepatitis A and Hepatitis B virus antigens (medicinal product)
- Vaccine product containing only Bordetella pertussis and Clostridium tetani and Corynebacterium diphtheriae antigens (medicinal product)

Has product characteristic and Has ingredient characteristic attribute values should be added as appropriate.

Example of Has product characteristic:

- Adult vaccine product containing only Hepatitis A virus antigen (medicinal product)
- Pediatric vaccine product containing only Hepatitis A virus antigen (medicinal product)
- Adult vaccine product containing only acellular Bordetella pertussis and Clostridium tetani toxoid and Corynebacterium diphtheriae toxoid antigens (medicinal product)

Example of Has ingredient characteristic:

- Vaccine product containing only Clostridium tetani and low dose Corynebacterium diphtheriae antigens (medicinal product)
- Vaccine product containing only Clostridium tetani and low dose Corynebacterium diphtheriae and inactivated Human poliovirus antigens (medicinal product)
Use the following pattern for the PT; align terming and case significance with the PT for the concept that is selected as the attribute value for the 127489000 |Has active ingredient (attribute)|. For multiple ingredient vaccine products, the active ingredients must be listed in alphabetical order, separated by the word "and", and the word "antigen" will be omitted. For concepts where all active ingredients are virus, the word "virus" may be omitted and added before "antigens".

- `<Active ingredient PT> only vaccine product`
- `<Active ingredient PT> and <Active ingredient PT> antigen only vaccine product`
- `<Active ingredient PT> and <Active ingredient PT> and <Active ingredient PT> antigen only vaccine product`

Example:

- Hepatitis B virus antigen only vaccine product
- Hepatitis A and Hepatitis B virus antigens only vaccine product
- Bordetella pertussis and Clostridium tetani and Corynebacterium diphtheriae antigens only vaccine product

Has product characteristic and Has ingredient characteristic attribute values should be added as appropriate.

Example of Has product characteristic:

- Hepatitis A virus antigen only adult vaccine product
- Hepatitis A virus antigen only pediatric vaccine product
- Adult acellular Bordetella pertussis and Clostridium tetani toxoid and Corynebacterium diphtheriae toxoid antigens only vaccine product

Example of Has ingredient characteristic:

- Clostridium tetani and low dose Corynebacterium diphtheriae antigens only vaccine product
- Clostridium tetani and low dose Corynebacterium diphtheriae and inactivated Human poliovirus antigens only vaccine product
Synonyms

Synonyms matching the FSN are not required.

Synonyms corresponding to the disorder that is the target of the vaccine are allowed. For multiple ingredient vaccine products, the disorders must be listed in alphabetical order and separated by the word "and". Note that these are not true synonyms; they may be updated and identified as "near-synonym" descriptions when that functionality becomes available.

Example:

- Hepatitis B vaccine
- Hepatitis A and Hepatitis B vaccine
- Diphtheria and pertussis and tetanus vaccine

Has product characteristic and Has ingredient characteristic attribute values should be added as appropriate.

Example of Has product characteristic:

- Hepatitis A adult vaccine
- Hepatitis A pediatric vaccine
- Diphtheria toxoid and acellular pertussis and tetanus toxoid adult vaccine

Example of Has ingredient characteristic:

- Low dose diphtheria and tetanus vaccine
- Low dose diphtheria and inactivated poliomyelitis and tetanus vaccine

Synonyms representing abbreviations for product (e.g. MMR, DTaP) will not be included in the International Release due to lack of internationally accepted reference source.

Exemplars

The following illustrates the **stated** view for 871822003 |Vaccine product containing only Hepatitis B virus antigen (medicinal product)|:

The following illustrates the **inferred** view for 871822003 |Vaccine product containing only Hepatitis B virus antigen (medicinal product)|:
The following illustrates the **stated** view for 871803007 [Vaccine product containing only Hepatitis A and Hepatitis B virus antigens (medicinal product)]:

The following illustrates the **inferred** view for 871803007 [Vaccine product containing only Hepatitis A and Hepatitis B virus antigens (medicinal product)]:
The following illustrates the **stated** view for 1991000221106 [Vaccine product containing only Human papillomavirus 16 and 18 antigens (medicinal product)]:

The following illustrates the **inferred** view for 1991000221106 [Vaccine product containing only Human papillomavirus 16 and 18 antigens (medicinal product)]:
The following illustrates the **stated** view for 865997008 |Adult vaccine product containing only Hepatitis A virus antigen (medicinal product)| and illustrates the use of the "Has product characteristic" attribute:

The following illustrates the **inferred** view for 865997008 |Adult vaccine product containing only Hepatitis A virus antigen (medicinal product)| and illustrates the use of the "Has product characteristic" attribute:
The following illustrates the **stated** view for 871838009 |Vaccine product containing only Clostridium tetani and low dose Corynebacterium diphtheriae and inactivated Human poliovirus antigens (medicinal product)| and illustrates use of the "Has ingredient characteristic" attribute:

```
871838009
Vaccine product containing only Clostridium tetani and low dose Corynebacterium diphtheriae and inactivated Human poliovirus antigens (medicinal product)
```

The following illustrates the **inferred** view for 871838009 |Vaccine product containing only Clostridium tetani and low dose Corynebacterium diphtheriae and inactivated Human poliovirus antigens (medicinal product)| and illustrates use of the "Has ingredient characteristic" attribute:
Role in Medicinal Product

See subsections for details re: editorial guidelines related to modeling and terming for roles in the Medicinal Product hierarchy.

Role Hierarchy

Overview

766940004 [Role (role)] and its descendant concepts are an abstract representation of a high-level role for a product; the concepts are not intended to describe a detailed indication for use nor imply that use is appropriate in all clinical situations.

The role hierarchy is comprised of concepts required to model or group concepts in SNOMED CT; it is not a comprehensive hierarchy. Concepts not required for modeling in SNOMED CT may be deprecated if not needed as groupers.

Note: Editorial guidelines for other types of roles will be added when a need for such concepts is identified and implemented.

Modeling

<table>
<thead>
<tr>
<th>Stated parent concept</th>
<th>766940004 [Role (role)] or an appropriate descendant of 766940004 [Role (role)]</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Exceptions: none identified</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(role)</th>
</tr>
</thead>
</table>

<table>
<thead>
<tr>
<th>Definition status</th>
<th>9000000000000074008</th>
<th>Necessary but not sufficient concept definition status (core metadata concept)</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>Exceptions: none identified</td>
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</table>

<table>
<thead>
<tr>
<th>Attributes</th>
<th>None</th>
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</table>
Naming

FSN

Use the following naming pattern for the FSN where X is a role. Each concept should represent a single role.

- X role (role)

Example:

- Active immunity stimulant role (role)

Use the following pattern for the FSN where X is a therapeutic role. Each concept should represent a single therapeutic role.

- X therapeutic role (role)

Example:

- Analgesic therapeutic role (role)
- Antifungal therapeutic role (role)

Preferred Term

Use the following naming pattern for the FSN where X is a role; align naming and case sensitivity with the FSN.

- X role

Use the following pattern for the PT; align naming and case sensitivity with the FSN.

- X therapeutic role

Example:

- Analgesic therapeutic role
- Antifungal therapeutic role

Synonym

Synonyms matching the FSN are not required.

Exemplars

The following illustrates the stated and inferred view for 773839009 |Analgesic therapeutic role (role)|:

The following illustrates the stated and inferred view for 788020005 |Antifungal therapeutic role (role)|:
Groupers Based on Role

Overview

Grouper concepts based on role that are deemed to be clinically useful and that can be sufficiently defined may be included in the |Medicinal product| hierarchy.

Note: This section applies to grouper concepts representing a single role.

A high-level grouper concept supports the organization of the hierarchy based on therapeutic role:

- 763087004 |Medicinal product categorized by therapeutic role (product)|

Modeling

<table>
<thead>
<tr>
<th>Stated parent concept</th>
<th>763158003</th>
<th>Medicinal product (product)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Exceptions: none identified</td>
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</table>

<table>
<thead>
<tr>
<th>Semantic tag</th>
<th>(product)</th>
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</thead>
<tbody>
<tr>
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<table>
<thead>
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<th>Definition status</th>
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<tbody>
<tr>
<td>Exceptions: none identified</td>
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<td></td>
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</tbody>
</table>

| Attribute: Plays role | Range: <766940004 | Role (role)| |
|-----------------------|------------------|-------------|
| Exceptions: none identified |

- Cardinality: 0..*
  - NOTE: While the allowed range is broader, the |Medicinal product| grouper concepts based on role should have one and only one |Plays role| attribute.
  - Exceptions: none identified

Naming Guidelines

<table>
<thead>
<tr>
<th>FSN</th>
<th>Use the following pattern for the FSN; align naming and case significance with the FSN for the concept that is selected as the attribute value, excluding the semantic tag and the words &quot;therapeutic role&quot;.</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>• Medicinal product acting as &lt;Therapeutic role FSN&gt; agent (product)</td>
</tr>
<tr>
<td></td>
<td>Example:</td>
</tr>
<tr>
<td></td>
<td>• Medicinal product acting as analgesic agent (product)</td>
</tr>
<tr>
<td></td>
<td>• Medicinal product acting as antacid agent (product)</td>
</tr>
<tr>
<td></td>
<td>• Medicinal product acting as antiglaucoma agent (product)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Preferred Term</th>
<th>Use the following pattern for the PT; align naming and case significance with the PT for the concept that is selected as the attribute value, excluding the words &quot;therapeutic role&quot;.</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt;Therapeutic role PT&gt; agent</td>
<td></td>
</tr>
<tr>
<td>Example:</td>
<td></td>
</tr>
<tr>
<td>• Analgesic agent</td>
<td></td>
</tr>
<tr>
<td>• Antacid agent</td>
<td></td>
</tr>
<tr>
<td>• Antiglaucoma agent</td>
<td></td>
</tr>
</tbody>
</table>

| Synonyms | Synonyms matching the FSN are not required. |
Exemplars
The following illustrates the **stated** view for grouper concept 788023007 |Medicinal product acting as antifungal agent (product)|:

The following illustrates the **inferred** view for grouper concept 788023007 |Medicinal product acting as antifungal agent (product)|:

The following illustrates the **stated** view for grouper concept 53009005 |Medicinal product acting as analgesic agent (product)|:

The following illustrates the **inferred** view for grouper concept 53009005 |Medicinal product acting as analgesic agent (product)|:
Groupers Based on Role plus Structure

Overview

Grouper concepts based on role plus structure that are deemed to be clinically useful and that can be sufficiently defined may be included in the |Medicinal product| hierarchy.

Note: This section applies to grouper concepts representing a single role plus a single structure.

Modeling

| Attribute: Has active ingredient | Range: <=105590001|Substance (substance) |
|----------------------------------|------------------|
|                                  | NOTE: While the allowed range is broader, the |Medicinal product| grouper concepts based on role plus structure should only use primitive grouer concepts that are descendants of 312413002 |Substance categorized by structure (substance)| as attribute values. |
|                                  | Exceptions: none identified |
|                                  | Cardinality: 0..* |
|                                  | NOTE: While the allowed range is broader, the |Medicinal product| grouper concepts based on role plus structure should have one and only one |Has active ingredient| attribute. |
|                                  | Exceptions: none identified |
### Attribute: Plays role

- **Range:** `<766940004 |Role (role)|`
  - Exceptions: none identified

- **Cardinality:** 0..*
  - NOTE: While the allowed range is broader, the |Medicinal product| grouper concepts based on role plus structure should have one and only one |Plays role| attribute.
  - Exceptions: none identified

### Naming Guidelines

**FSN**

Use the following pattern for the FSN; align naming and case significance with the FSN for the concept that is selected as the attribute value, excluding the semantic tag and the words "therapeutic role".

- Medicinal product containing `<Active ingredient PT>` and acting as `<Therapeutic role FSN>` agent (product)

**Example:**

- Medicinal product containing tricyclic and acting as antidepressant agent (product)
- Medicinal product containing azole and acting as antifungal agent (product)

**Preferred Term**

Use the following pattern for the PT; align naming and case significance with the PT for the concept that is selected as the attribute value, excluding the words "therapeutic role".

- `<Active ingredient PT>` `<Therapeutic role PT>` agent

**Example:**

- Tricyclic antidepressant agent
- Azole antifungal agent

**Synonyms**

Synonyms matching the FSN are not required.

### Exemplars

The following illustrates the **stated** view for grouper concept 350195009 |Medicinal product containing azole and acting as antifungal agent (product)|:

```
+-----------------+-----------------+-----------------+-----------------+
| 350195009       | Medicinal product containing azole and acting as antifungal agent (product) |
|-----------------+-----------------+-----------------+-----------------|
| ➔                |                 | 769939001       | Plays role (attribute) |
|                 |                 | 766940004       | Role (role) |
|                 |                 | 788020005       | Antifungal therapeutic role (role) |
|                 |                 | 157489000       | Has active ingredient (attribute) |
|                 |                 | 300939003       | Azole (substance) |
|                 | 769168003       | Medicinal product (product) |

The following illustrates the **inferred** view for grouper concept 350195009 |Medicinal product containing azole and acting as antifungal agent (product)|:
The following illustrates the **stated** view for grouper concept 33219003 |Medicinal product containing tricyclic compound and acting as antidepressant agent (product)|:

The following illustrates the **inferred** view for grouper concept 33219003 |Medicinal product containing tricyclic compound and acting as antidepressant agent (product)|:
Modeling Association between Role and Product

Overview
Associations between role and product in the Medicinal product hierarchy are primarily created to support modeling in other hierarchies; however, organization of content or facilitation of maintenance activities may also be reasons for creation of associations.

Modeling
Associations between therapeutic role and concepts in the Medicinal product hierarchy shall be modeled by creation of an additional non-defining axiom.

The additional axiom should be added to the most general possible concept with one of the following semantic tags:

- (medicinal product)
- (medicinal product form)
- (clinical drug)

A separate additional axiom must be created for each therapeutic role.

| Stated parent concept for additional axiom | 763158003|Medicinal product (product) |
|------------------------------------------|------------------------------------------|
| Exceptions: none identified |
| Note: Addition of the additional axiom does not impact the editorial guidelines for modeling or terming for the (medicinal product), (medicinal product form), or (clinical drug) concept as described in preceding sections of this document. |

| Definition status for additional axiom | 900000000000074008 |Not sufficiently defined by necessary conditions definition status (core metadata concept)| |
|---------------------------------------|------------------------------------------|
| Exceptions: none identified |

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Attribute: Plays role
- Range: <=766940004 |Role (role)|
  - Exceptions: none identified
- Cardinality: 0..*
  - NOTE: While the allowed range is broader, the additional axiom should have one and only one |Plays role| attribute.
  - Exceptions: none identified

Exemplars
The following illustrates the stated view for concept 40589005 |Product containing amitriptyline (medicinal product)|:

The following illustrates the inferred view for concept 40589005 |Product containing amitriptyline (medicinal product)|:

The following illustrates the stated view for concept 431023006 |Product containing dapsone in oral dose form (medicinal product form)|:
The following illustrates the **inferred** view for concept 431023006 |Product containing dapsone in oral dose form (medicinal product form)|:
Patch Test Product

411126008 |Patch test product (product)| and its descendants will be retained "as is" until use cases and/or detailed requirements are known.

Requests for addition of new concepts will be rejected. Requests for modification of existing concepts will be evaluated on a case-by-case basis.

Sterile Maggots Product

410969008 |Sterile maggots (product)| and its descendants will be retained as a primitive subhierarchy until use cases and/or detailed requirements are known.

Requests for addition of new concepts or modification of existing concepts will be evaluated on a case-by-case basis.

Appendix A - Cumulative Summary of Changes for Pharmaceutical and Biologic Product Editorial Guide

<table>
<thead>
<tr>
<th>Date</th>
<th>Comments</th>
<th>Effective date</th>
</tr>
</thead>
<tbody>
<tr>
<td>2021-Jul</td>
<td>Updated to reflect:&lt;br&gt;  • Concrete domain deployment&lt;br&gt;  • New attributes for Has target population and Has qualitative strength&lt;br&gt;  • General updates including clarifications and refinements of editorial guidelines</td>
<td>2021-Jul</td>
</tr>
<tr>
<td>2021-Jan</td>
<td>Revisions to support making Pharmaceutical and Biologic Product SNOMED CT Editorial Guide available via SNOMED International Document Library.</td>
<td>2021-Jan</td>
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