



## James Read Memorial Lecture SNOMED CT - A Canadian Clinical Perspective

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### Overview

- Overview of SNOMED CT in Canada
- SNOMED CT implementation in Hospitals – Challenges and Solutions:
  - Past: “stealth mode”
  - Present: “building into daily care”
- The (near) Future of SNOMED CT in Canada:
  - Provincial clinical standardization
  - Evidence-based content distribution
  - Iterative quality improvement

## Working as a CMIO = Parenting



## Canada Health Infoway and SNOMED CT



- Advocates for SNOMED CT use
- Maintains SNOMED CT Canadian Edition (EN/FR)
- Provides access to SNOMED CT (via license)
- Supports and educates users and developers
- Manages requests for content changes/updates (RFC's)
- Oversees development of Canadian Subsets
  - Immunization, Communicable Disease, ePrescribing, Primary Care

## SNOMED CT: Challenges for Canadian Hospitals

- SNOMED CT not the default terminology provided by vendors
- SNOMED CT Edition / National Extension confusion
- Poor organizational/clinical leadership appreciation re: benefits of SNOMED CT for clinical standardization, interoperability, decision support
- Poor vendor support for effective searches/clinician workflows:
  - Additional multi-disciplinary expertise required for custom interfaces, documentation templates with selected/validated concept subsets
  - User adoption challenges → overreliance on free text entries (50%)
- User frustration: volume of concepts, missing synonyms, redundant terms

“...there are very few known SNOMED CT implementations in (hospital) clinical care settings.”

Liu J, Lane K, Veillette C et al. Addressing SNOMED CT Implementation Challenges through Multi-Disciplinary Collaboration. *Stud Health Tech Inf* 2010; 981-985.

Lee D, Cornet R, Lau F, de Keizer N et al. A survey of SNOMED CT implementations. *J Biomed Inf* 2013; 46: 87-96.

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Community academic hospital affiliated with the University of Toronto

**Catchment area:** > 400,000

**Three Sites:** General, Branson, Seniors' Health

**Beds:** 426 acute care  
192 long-term care

**Volumes per year:**

- 124,000 ED visits
- 31,000 inpatient cases
- 214,000 outpatient cases
- 5,800 births

**HIMSS Analytics**  
**STAGE 6**  
**2011**

**HIMSS**  
Davies Award  
NICHOLAS E. DAVIES  
**2016**

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## What is eCare?

Advanced Hospital Information System (HIS), with CPOE and electronic documentation

+

Standardization on Evidence-Based Care

+

Safe Prescribing and e-Medication Management

+

Clinical Decision Support (Static and Dynamic)



Kickoff: **2007**

Phased Implementation:  
**2008-2015**

Hospital-wide: **2015**

## Goals of the eCare Project



- **Implement advanced HIS to *improve patient outcomes*:**
  - **Quality and safety of patient care**
  - **Enable Clinical & Business Intelligence for better decisions**
- **Embrace culture of standardized, evidence-based care**
  - **Build evidence and best practice into optimized workflows**
  - **Make it “easy to do the right thing”**
- **SHARED VISION = “by clinicians, for clinicians”**
  - **100% clinician adoption via comprehensive engagement**
  - **Team-based interprofessional approach/workflows**

## Success Factors for SNOMED-CT Clinical Implementation

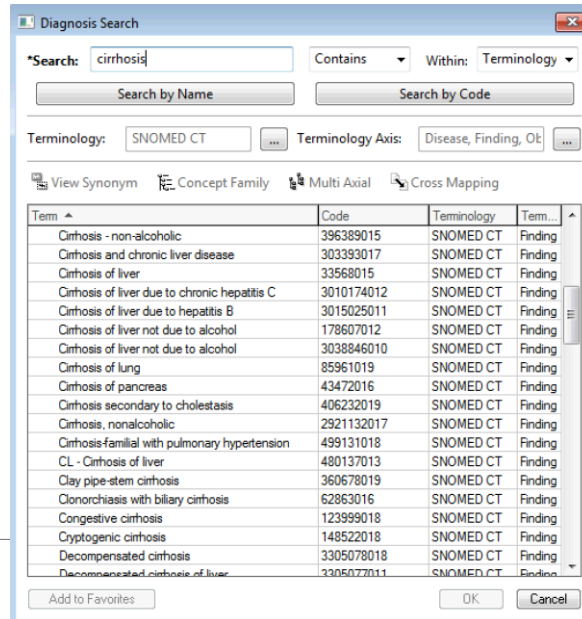
1. **Simplicity:** hide the complexity, “Google search”
2. **Clinician engagement:** “understand the why”
3. **Demonstrate value:** clinically relevant
4. **Reference sites:** “have a mentor”
5. **Training:** only helpful to a point (see #1)
6. **Vendor assistance:** system must support efficient workflow, accurate concept selection

## Helping MD’s “Understand the Why”

- Clinically relevant, granular, comprehensive, flexible terminology
- Designed for direct use by the clinician (vs post-coding by analysts):
  - Highest accuracy and clinical utility
  - Cross-mapping allows better resource intensity weighting = better hospital funding
- Change the channel – big-picture workflow instead of click-counting:
  - Problem lists that automatically populate every consult/progress/discharge note
  - Problem lists carry between hospital visits
  - Physician handover list automatically supports active problems
- Driving clinical decision support:
  - Real-time: suggestion of order sets, disease-drug interactions
  - Longitudinal: screening recommendations (e.g. polyps based on family/personal hx)
  - Population: clinically accurate diagnoses managed across facility, region, province
- Quality Improvement, Research, Resource Management:
  - SNOMED CT coded data drives all three activities (better data accuracy/availability)

## Problem List: Clinical Adoption Challenges

- Too many terms to review and select (e.g. “hypertension”)
- Average 12 seconds per diagnosis
- Viewed as “clerical” (though already done on paper)  
→ *not in workflow*
- **< 1% adoption**



11



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## Success Factors for SNOMED-CT Clinical Implementation

- ~~1. **Simplicity:** hide the complexity, “Google search”~~
- ~~2. **Clinician engagement:** “understand the why”~~
- ~~3. **Demonstrate value:** clinically relevant, “big picture”~~
- ~~4. **Reference sites:** “have a mentor”~~
- ~~5. **Training:** only helpful to appoint (see #1)~~
- ~~6. **Vendor assistance:** system must support efficient workflow, accurate concept selection~~

12



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# Starting in Stealth Mode

**“Make it easy to do the right thing”**

Use SNOMED CT in daily clinical workflow ...  
without realizing it



## Pneumonia Admission Order Set: Evidence-Based Empiric Antibiotic Treatment Selection

**Guidelines**

- Please refer to NYGH Antimicrobial Guideline Handbook, and select appropriate empiric antimicrobial regimens during first 24 hours. Use oral option when possible unless NPO or vomiting
- Antibiotics should be administered as soon as possible (within 4-6 hours) after the diagnosis of CAP has been made in ER. If patient has taken an antibiotic within the last 3 months for any reason, select an antibiotic from an ALTERNATE class
- EARLY SWITCH FROM PARENTERAL TO ORAL ANTIMICROBIAL THERAPY: should be considered followed by discharge for eligible patients (e.g. hemodynamically stable, improving clinically, normal GI tract, and able to ingest medications)
- For suspected aspiration OR Healthcare Associated Pneumonia (HCAP): Use the following modules instead of the antimicrobial options below

Pneumonia With Suspected Aspiration (Adult) (Modul...)

Pneumonia: Nosocomial/Health Care Associated Pne...

**Antibiotic Regimens**

- First-line treatment: Beta-lactam (ceftriaxone, or if taking PO, amoxicillin-clavulanic acid). Addition of a macrolide is indicated for patients with severe illness, positive urine antigen, or suspected Legionellosis (See Atypical Coverage: Macrolide section below)
- In patients with a true beta-lactam allergy (eq. anaphylaxis, angioedema, or bronchospasm): Respiratory fluoroquinolone alone
- DURATION OF THERAPY: 5-7 days of therapy for hospitalized patients not in ICU

**First Line Treatment: Beta-Lactam**

<input checked="" type="checkbox"/>	cefTRIAxone (Rocephin inj)	1000 mg, Inj, IV, q24h-ATC, NOW
<input checked="" type="checkbox"/>	amoxicillin-clavulanate (Clavulin F 875-125)	875 mg, PO, q12h, NOW
<input checked="" type="checkbox"/>	amoxicillin-clavulanate (Clavulin F 500-125)	500 mg, Tab, PO, q12h, NOW, Suggested dose for creatinine clearance 10-30 ml...

**Atypical Coverage: Macrolide**

- A macrolide for atypical coverage is indicated in patients with severe illness, positive urine antigen, or suspected Legionellosis

<input checked="" type="checkbox"/>	azithromycin (Zithromax inj)	500 mg, Inj, IV, q24h-ATC, for 5 days, NOW
<input checked="" type="checkbox"/>	azithromycin (Zithromax)	500 mg, Tab, PO, q24h-ATC, NOW

**Beta-lactam Allergy: Quinolone**

- A respiratory fluoroquinolone is indicated in patients with a true beta-lactam allergy (eq. anaphylaxis, angioedema, or bronchospasm)

<input checked="" type="checkbox"/>	Moxifloxacin (IV or PO) is indicated for low to intermediate risk patients	
<input checked="" type="checkbox"/>	moxifloxacin (Avelox)	400 mg, Tab, PO, q24h-ATC, NOW
<input checked="" type="checkbox"/>	moxifloxacin (Avelox IV.)	400 mg, Inj, IV, q24h-ATC, NOW

For renally impaired patients, no dose adjustment is required with moxifloxacin

**Suspected Pseudomonas**

- First-line treatment for SUSPECTED/PROVEN P.AERUGINOSA: piperacillin-tazobactam +/- azithromycin
- For patient with true beta-lactam allergy: Meropenem +/- azithromycin. Meropenem is associated with low cross-reactivity among those with a beta-lactam allergy.

**Reminder**

For patients who have atrial fibrillation and are at high or intermediate risk for stroke, use oral anticoagulation with apixaban, dabigatran, rivaroxaban, or warfarin.

For those who are unsuitable for warfarin therapy, use a direct thrombin inhibitor or factor Xa inhibitor.

For patients who have atrial fibrillation of  $\geq 48$  hours' duration or of unknown duration and who are undergoing cardioversion, use warfarin, an LMWH, apixaban, rivaroxaban, or dabigatran for at least 3 weeks prior to and 4 weeks after non-TEE-guided cardioversion.

For those who have atrial fibrillation of  $< 48$  hours' duration and are undergoing cardioversion, use heparin (either IV UFH or an LMWH), a direct thrombin inhibitor, or factor Xa inhibitor if the patient is not already anticoagulated.

For patients with atrial fibrillation who have a creatinine clearance  $< 25$  mL/minute, do not use apixaban.

For patients with atrial fibrillation who have a creatinine clearance  $< 15$  mL/minute, do not use rivaroxaban.

Abbreviations | Guidelines

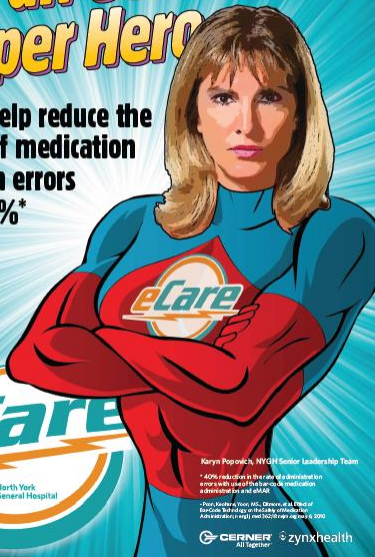
**Rationale**

The following table summarizes meta-analyses related to this topic:

Study	Comparison	
Giugliano et al (ENGAGE AF-TIMI 48, 2014)	High-dose (60 mg once daily) edoxaban vs Low-dose (30 mg once daily) edoxaban vs Warfarin	In patients with atrial fibrillation who have moderate-high stroke risk: <ul style="list-style-type: none"> <li>High-dose edoxaban decreases stroke (ischemic or hemorrhagic) during treatment</li> <li>There is no significant between-group difference in stroke during treatment</li> <li>There is no significant between-group difference in a combined outcome of stroke or death</li> <li>Low-dose edoxaban increases the frequency of a combined outcome of stroke or death</li> <li>Both high-dose and low-dose edoxaban decrease hemorrhagic stroke and death</li> </ul>
Hajperin et al (ROCKET AF, 2014)	Rivaroxaban vs Warfarin	In patients with nonvalvular atrial fibrillation at moderate to high risk of stroke: <ul style="list-style-type: none"> <li>In patients <math>\geq 75</math> years of age:                             <ul style="list-style-type: none"> <li>Rivaroxaban increases the combined outcome of major or clinically significant bleeding</li> <li>There is no significant between-group difference in a combined outcome of stroke or death</li> </ul> </li> <li>In patients <math>&lt; 75</math> years of age:                             <ul style="list-style-type: none"> <li>There is no significant between-group difference in a combined outcome of stroke or death</li> <li>There is no significant between-group difference in a combined outcome of stroke or death</li> </ul> </li> </ul>
Lip et al (2014)	Apixaban vs Aspirin	Based on data from the AVERROES study and at a mean follow-up of 1.1 years, <ul style="list-style-type: none"> <li>In all patients, apixaban decreases ischemic stroke</li> <li>In female patients, apixaban decreases ischemic stroke</li> <li>In male patients, apixaban decreases ischemic stroke</li> <li>In all patients, there is no significant between-group difference in intracranial hemorrhage</li> <li>In female patients, there is no significant between-group difference in intracranial hemorrhage</li> <li>In male patients, there is no significant between-group difference in intracranial hemorrhage</li> </ul>
Hylek et al (2014)	Apixaban vs Warfarin	Based on data from the ARISTOTLE trial, in patients with atrial fibrillation, apixaban decreases stroke (ischemic or hemorrhagic) during treatment
Antang et al (2013)	Warfarin vs Alternative anticoagulant (eg, direct thrombin inhibitors, factor Xa inhibitors, aspirin, clopidogrel)	In patients with atrial fibrillation, there is no significant between-group difference in stroke (ischemic or hemorrhagic) during treatment
Bruins Slot and Berge (CD008980, 2013)	Factor Xa inhibitors vs Vitamin K antagonists	In patients with atrial fibrillation or atrial flutter, factor Xa inhibitors reduce the risk of stroke, major bleedings, intracranial hemorrhages, and all-cause deaths.

## Be an eCare Super Hero

**and help reduce the rate of medication admin errors by 40%\***



Karyn Popovich, NYGH Senior Leadership Team


\* 40% reduction in the rate of medication admin errors with eCare (NYGH) compared to medication administered per the standard.

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## Be an eCare Super Hero

**and help save 21% of inpatients lives at risk from pneumonia\***



Dr. David Barish, NYGH Physician

\* 21% reduction in inpatient deaths from pneumonia using eCare (NYGH) compared to standard of care.

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## Driving Problem List from Order Sets

- >97% adoption of diagnosis-specific admission order sets
- Build quick-click context-specific diagnoses and comorbidities into ordering workflow
- **From <1% adoption to 15% adoption**

Discern: (2 of 2)

### Comorbidities?

Does this patient have any of the comorbidities below? You can select multiple options where applicable.

If none of these comorbidities apply, click 'OK' without making any selection.

PLEASE NOTE: Later, if you wish to add/modify comorbidities for this patient, go to the left-hand Menu and click "Diagnoses & Problems".

Add Problem(s) for:

- Atherosclerotic heart disease
- Chronic congestive heart failure
- Chronic left-sided congestive heart failure
- Chronic right-sided congestive heart failure
- Hypertension
- Increased lipid
- Smoking

Diagnosis (Problem) being Addressed this Visit

+ Add    Modify    Convert    Display: All

Clinical Dx	Date	Dx Type	Code
Acute congestive heart failure	2017-Oct-17	Working	18472010

## Ambulatory Synoptic Reporting Templates

- Built context-sensitive clickable SNOMED CT encoded diagnoses into documentation workflow: endoscopy, diabetes care, urology
- Required extensive clinician input, terminology expertise
- **From 15% adoption to 30% adoption**
- **Analytics for quality improvement – e.g. polypectomy rate by physician**

	Total cases	Polyp Seen	Polypectomy	Polyp Detection Rate	Polypectomy Rate
Surgeon A	179	78	71	43.6%	39.7%
Surgeon B	692	253	233	36.5%	33.7%
Surgeon C	480	123	113	25.6%	23.5%
Surgeon D	128	33	22	25.8%	17.2%
Surgeon E	167	36	23	21.5%	13.8%

# Real-Time Clinical Decision Support: Drug-Disease Interaction

Discern: (1 of 1)

## Potential Inappropriate Antipsychotic Use

Only prescribe quetiapine or low dose parenteral olanzapine for agitation if patient has Parkinson's Disease or Lewy Body Dementia.

### Alert Action

- Cancel Haldol
- Ignore

OK

**NORTH YORK GENERAL** Physician Scorecard - Medicine  
Reporting Period: FY1516  
Physician Group - Cardiology

#### Statistics for Physician Group - Cardiology (For Congestive Heart Failure)

Cases	Weighted Cases (ARIW)	Average Total LOS (days)	Average Acute LOS - (25th Percentile Target) - (days)	Conservable Days	ALC as % Total IP Days	Re-Admits within 30 days	Weekend Discharge %
491	653 (1.3)	6.6	6.0 (3.6)	926 (37%)	9%	55 (13.3%)	11%

#### Comparisons to Medicine Program (For Congestive Heart Failure)

Cases	Weighted Cases (ARIW)	Average Total LOS (days)	Average Acute LOS - (25th Percentile Target) - (days)	Conservable Days	ALC as % Total IP Days	Re-Admits within 30 days	Weekend Discharge %
926	1,358 (1.5)	7.6	6.8 (3.9)	1,834 (36%)	11%	107 (14.2%)	12%

**Physician Scorecards**

#### Physician Detail (For Congestive Heart Failure)

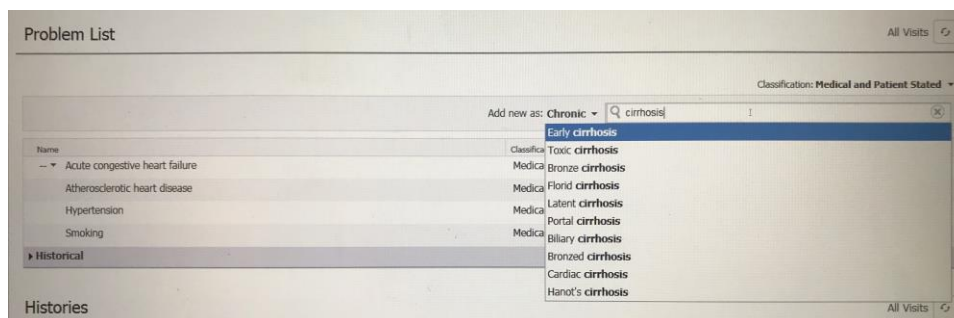
Physician	Cases	Weighted Cases (ARIW)	Average Total LOS (days)	Average Acute LOS (days)	Conservable Days	ALC as % Total IP Days	Re-Admits within 30 days	Weekend Discharge %
	66	81 (1.2)	6.9	5.7 (3.5)	132 (39%)	18%	8 (15.1%)	14%
	82	105 (1.3)	5.6	4.9 (3.3)	43 (17%)	11%	7 (10.0%)	11%
	76	99 (1.3)	7.6	6.5 (3.5)	177.7 (44%)	14%	8 (12.5%)	14%
	75	133 (1.8)	7.2	7.0 (3.9)	188 (41%)	3%	8 (12.7%)	15%
	75	84 (1.1)	6.2	5.3 (3.4)	133.4 (36%)	15%	11 (19.0%)	7%
	68	82 (1.2)	6.1	6.0 (3.6)	145 (38%)	1%	11 (19.6%)	13%
	65	86 (1.3)	6.9	6.4 (3.7)	130.7 (38%)	8%	6 (10.7%)	14%
	50	64 (1.3)	6.3	6.0 (3.9)	108 (36%)	5%	4 (8.7%)	2%

## Integrating SNOMED-CT Into Daily Physician Inpatient Documentation

- Vendor introduced new documentation software: builds easier-to-use problem list into improved clinical documentation workflow, SNOMED CT capable
- Simplicity, demonstrable value, clinician engagement barriers lowered
- Approach:
  - New clinically-focused problem list search algorithm and user interface:
    - Filtered top 10 choices that match search string, preferred terms and synonyms prioritized
    - Limited to concept types relevant to a problem list (e.g. finding, disorder, procedure...)
  - Pilot with 10 physicians, then specialty-by-specialty rollout
  - Problem list required to generate discharge summary
  - No free text diagnoses permitted, but free text comments allowed

## Integrating SNOMED-CT Into Daily Physician Inpatient Documentation

- Challenges:
  - “Top 10 match doesn’t always work”  
e.g. cirrhosis (unspecified), uncommon form of hypertension
  - Risk of miscoding – physicians may use “closest match”
  - No support for post co-ordination of terms (problems with laterality, etc)
- **100% adoption among pilot physicians**





## What Are the Results? Selected Outcomes from NYGH eCare



## HSMR:

- Reported from hospitals to CIHI annually
- Reported to public by CIHI annually
- GOAL: Reduce preventable inpatient deaths



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## Study: CPOE and Evidence-Based Order Sets

### Retrospective chart review:

- All patients discharged with a main diagnosis of Pneumonia or COPD
  - **Population #1:** Pre-CPOE (Jan-Sep 2010) n = 520
  - **Population #2:** Post-CPOE (Jan-Sep 2011) n = 511
  - Groups similar in age, gender distribution
  - Corrections: "Probability of Death", critical care admission

### Primary Hypothesis:

- Use of CPOE is associated with reduction in adjusted mortality vs traditional paper processes

### Secondary Hypothesis:

- Use of CPOE with a matching evidence-based admission order set is associated with reduction in adjusted mortality vs use of any order set

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## Results: CPOE vs Paper

Outcome	Odds Ratio	Confidence Interval	p-value
Death	<b>0.574</b>	0.391 – 0.843	<b>0.005</b>
Death adj for Probability of Death	<b>0.571</b>	0.383 – 0.852	<b>0.006</b>
Death adj for Probability of Death and CrCU Admission	<b>0.547</b>	0.360 – 0.830	<b>0.005</b>
30-Day Readmission	0.835	0.573 – 1.210	0.345
30-Day Readmission adj for Probability of Death and CrCU Admission	0.837	0.562 – 1.250	0.380

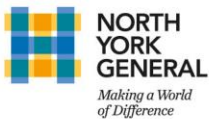
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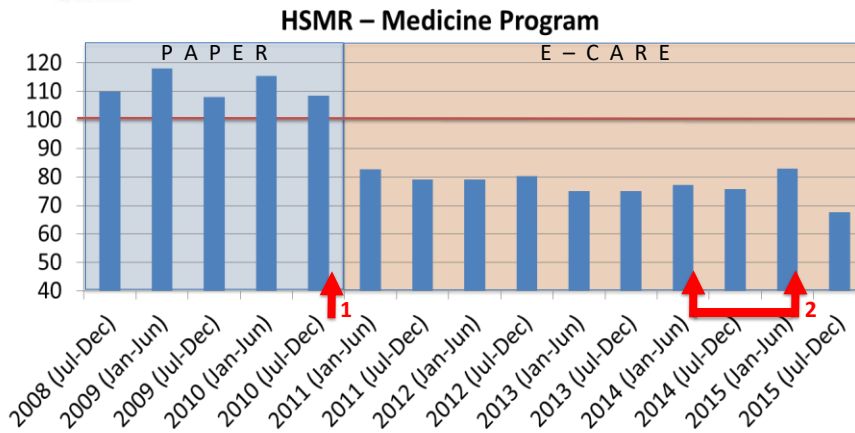
## Results: Evidence-Based Order Set Selection

Order Set	Outcome	Odds Ratio	Confidence Interval	p-value
Diagnosis-appropriate	Death	<b>0.48</b>	0.26 – 0.90	<b>0.022</b>
Diagnosis-appropriate	Death adj for Probability of Death and CrCU Admission	<b>0.44</b>	0.21 – 0.90	<b>0.024</b>
Diagnosis-appropriate	30-Day Readmission	1.35	0.75 – 2.38	0.30
Close to diagnosis	Death	1.47	0.71 – 3.01	0.30
Close to diagnosis	Death adj for Probability of Death and CrCU Admission	1.82	0.78 – 4.23	0.16
Any order set	Death	0.55	0.12 – 2.54	0.44
Any order set	30-Day Readmission	1.53	0.19 – 11.92	0.69

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## Inpatient Preventable Mortality: Trended Format

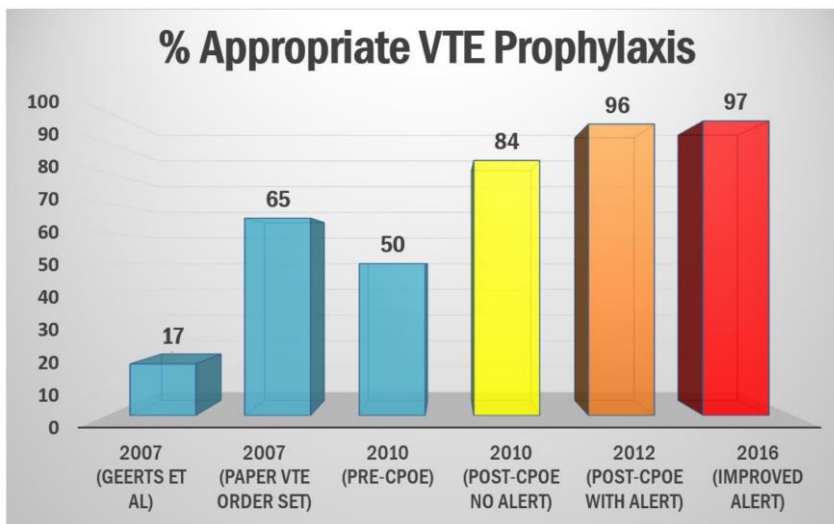


- 1 – eCare Phase 2 Implementation (CPOE, order sets, electronic med management)
- 2 – Quality Based Procedure (QBP) implementation – phased, over 1 year

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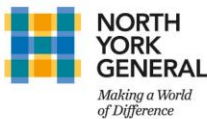


## Making Quality Stick: VTE Prophylaxis



Case: Venous Thromboembolism Prophylaxis

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# Summary of eCare Clinical Benefits

- **100%** clinician adoption, with over **80%** of clinicians “satisfied” or “very satisfied” with the system
- Medication reconciliation improved from **8%** to **>85%** of our medical patient population
- Medication turnaround time for STAT antibiotics improved by **83%** (291→50 mins)
- Over **11,000** potential medication administration errors averted (patient mismatch averted through closed-loop medication scanning)
- Appropriate prophylaxis against venous thromboembolism (VTE) increased from **50%** of inpatients to **>97%** of inpatients, with a corresponding **39%** reduction in VTE
- Order set usage on patient admission to hospital increased from **36.5%** (paper) to **>97%** (CPOE), even though use not mandatory
- Mortality from pneumonia and COPD exacerbation was reduced by **45%** using CPOE vs paper orders, and by **56%** using CPOE with a correctly-matched evidence-based order set

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## eCare ROI Calculation

Canadian cost of adverse nosocomial events:

- Cost per medication error: \$402 to \$632 (median **\$517 CDN**)
- Cost per nosocomial adverse drug event: **\$4,028 CDN**
- Cost per case of nosocomial VTE: **\$24,411** **\$36,047 CDN**
- Cost per case of nosocomial pneumonia: **\$17,000** **\$19,000 CDN**

Nosocomial Event	Cost Savings (\$CDN) Dec 2015
Medication Errors	\$30,428
Adverse Drug Events/Discrepancy	\$31,062,113
VTE prevention	\$1,029,169
Prevented recurrences of <i>C.difficile</i>	\$293,376
<b>TOTAL COST AVERTED</b>	<b>\$38,115,113</b>

Cost of >150 lives saved in 5 years from pneumonia and COPD exacerbation:

**PRICELESS**

*In Acute Care – CPSI July 2012*

→ Net savings over 5 years: **\$1.2 million**





## Utilizing SNOMED CT to Improve Quality of Hospital Care Nationally

### The Provincial and National Challenge

**Our results  
are not  
typical !**



- Most Canadian hospitals are not effective at integrating current evidence, standardized data into clinical workflows
- Reasons: leadership, resources/expertise (build/update), technology focus, poor application of standards
- Duplicate work: NYGH: **4.5 FTE**, **850** order sets, **~350** order sets updated annually

## IMPORTANCE OF CLINICAL CONTENT

Venue	Annual Savings: Efficiency from Automation	Annual Savings: Evidence-Based Care, Clin. Decision Support	Total Savings
Ambulatory	\$1.6 B (15%)	\$ 9.0 B (85%)	\$10.6 B
Inpatient	\$8.3 B (26%)	\$22.9 B (74%)	\$31.2 B
<b>TOTAL</b>	<b>\$9.9 B (24%)</b>	<b>\$31.9 B (76%)</b>	<b>\$41.8 B</b>

*Hillestad et al, Health Affairs 2005*

76% of the savings are from **better clinical decisions**, not efficiencies from automation

**“The most expensive tool used in medicine is the doctor’s pen”**



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## Ontario HIS Benefits and Adoption Team (HISBAT)

- Led by North York General Hospital (HIMSS 6) and Ontario Shores Centre for Mental Health Sciences (HIMSS 7), both Davies Enterprise Award winners
- Provided at no cost to Ontario hospitals (80% in need of assistance)
- Peer-to-peer knowledge sharing, mentorship of HIS project teams through on-site visits:
  - Governance, implementation, clinician engagement, standardized clinical content including order sets, terminology (SNOMED CT)
- First 9 months – 50+ hospitals assisted

36



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Ontario Shores  
Centre for Mental Health Sciences



Ontario



**HIS Adoption and  
Benefits Team**

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# Knowledge Translation for Hospital Information Systems: The Issues



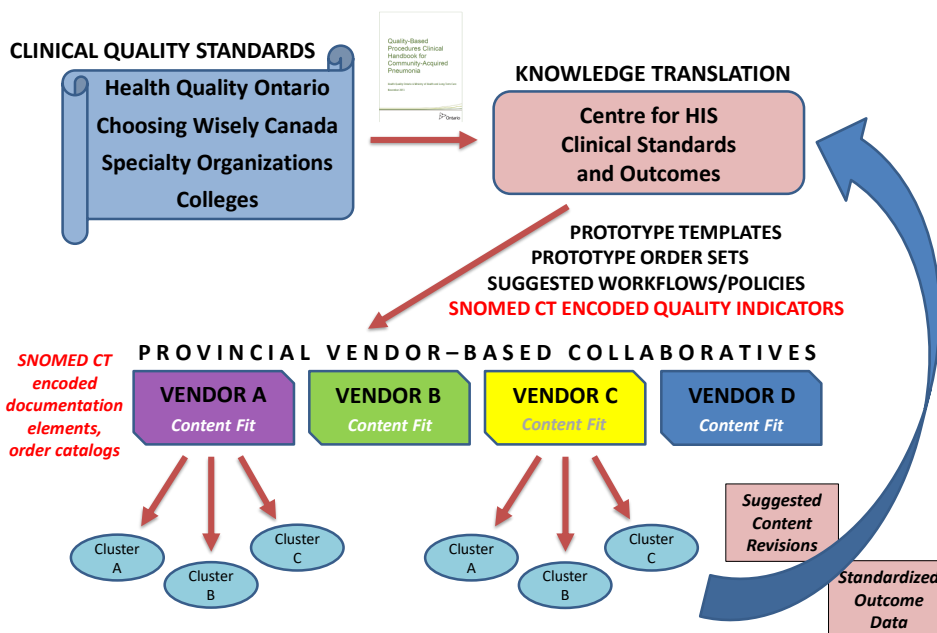
- Current provincial program attempts to use paper-based order sets to standardize quality implementation
- Some items are measurable against physician orders, but some are not (order may be a surrogate measure)
- Translation required from paper content to HIS: heterogeneous, time/resource intensive
- Elements in hospital information systems are free-text, not standardized

**Chemistry**

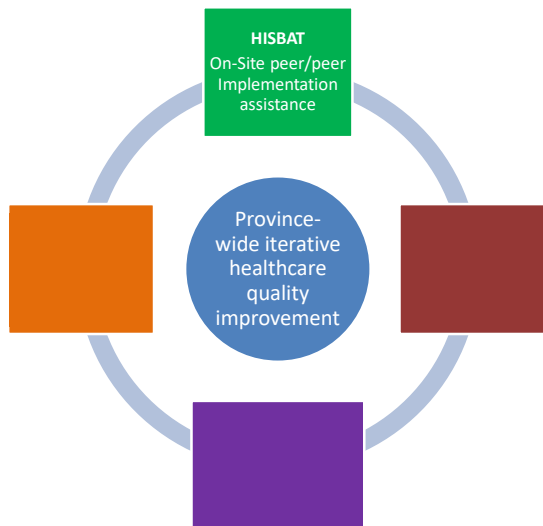
- Electrolytes **QBP**
- Creatinine **QPP**

Patients who do not have up-to-date influenza (annual) or pneumococcal vaccinations should either be vaccinated before discharge or referred for vaccination afterward, unless contraindications are present.

## Provincial Schematic: Clinical Standardization



## Provincial eHealth Clinical Quality Activities



## SNOMED CT: the Canadian Clinical/Hospital Journey So Far



### The Past:

- Misunderstanding/resistance



### The Present:

- Understanding
- Stealth use → Regular use
- Clinical benefits tangible



### The (near) future:

- Peer-to-peer mentorship
- Standardization of HIS “building blocks”
- Centralized, quality-focused clinical content dev.
- Closed loop: system-level analysis/improvement
- Translatable approach for publicly-funded jurisdictions





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of Difference*

**THANK YOU!**

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