Use of Clinic/Physician Office (C/PO) Electronic Health **Records (EHRs) to Improve Cancer Surveillance Quality**, **Reduce Costs and Advance** ePublic Health



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Background: Hospital Reporting

- Central cancer registries (CCRs) have traditionally relied on hospitals to abstract and submit new cancer cases to the CCR
 - Advantage
 - Relatively small number of hospital systems
 - Except for low-volume facilities, have trained registry staff that review medical records and prepare abstracts

Background: Changes in Reporting

- Changes in medical practice and health care delivery have increased the incidence of cancers being diagnosed/treated outside a hospital setting
 - For example, within Missouri:
 - inpatient & outpatient hospital
 - pathology laboratories
 - ambulatory surgical centers
 - free-standing cancer clinics and treatment centers
 - skilled nursing facilities
 - intermediate care facilities
 - residential care facilities I and II
 - physician offices (if not reported by another source)
 - Note: Collecting non-inpatient cases required expansion of cancer reporting laws

Background: Changes in Reporting (Cont'd)

- Complete surveillance of some cancers may be particularly affected by diagnosis/ treatment outside of the hospital setting, *e.g.*,
 - Leukemia
 - Lymphoma
 - Melanoma skin cancer
 - Prostate cancer
 - in situ & localized Breast, Cervical, and Colorectal cancers

Background: C/PO Reporting

- Cases treated entirely within a physician office setting present a potential challenge for CCRs
 - Relatively large number of C/POs
 - Lack of trained tumor registrars to prepare abstracts

Background: C/PO Reporting (Cont'd)

- But EHR systems provide a possibility of automating the collection of detailed data
 - The majority of NPCR-funded CCRs only collect 1st course treatment, summary treatment information, and no co-morbidities
 - This limits the ability to evaluate outcomes and comparative effectiveness research
 - Access to EHR data can potentially provide treatment and co-morbidities detail not present in standard cancer abstracts
 - Challenges: case selection, storage

Background: Current Sources

- 120 dermatologists report to MCR
 - 19 (16%) electronically via CDC's Web Plus[™]
 - 101 (84%) via paper form
- MCR has been receiving ePath reports (non-CDA based) via PHIN-MS for several years
- Free-Standing Radiation Facilities report prostate via CDC's Web Plus
- Ambulatory Surgical Centers report via paper

Background: Current Sources (Cont'd)

Hospitals

- Non-Low Volume Facilities report electronically
- Low Volume Facilities (including Critical Access Hospitals [CAHs])
 - Currently, MCR or a contractor receives copies of medical records to abstract
 - Efficiency can potentially be increased by electronic reporting
- MCR-ARC is one of two CCRs that participated in an ARRA-funded pilot project to improve cancer reporting by importing real-time data directly from EHRs to CCR

Methods: Identifying Partners

- To increase case completeness by obtaining previously unreported cases and treatment information from EHRs, we:
 - Partnered with the Missouri Health Information Technology (MO HIT) Assistance Center to identify potential:
 - Clinic/physician offices (C/POs)
 - Critical access hospitals (CAHs)

Methods: Identifying Partners (Cont'd)

- Conducted site visits
 - Recruited 8 participants
 - 6 CAHs, 2 C/POs
 - Focus primarily on the 2 C/POs
- Identified and collaborated with:
 - Facility EHR vendors
 - CDC software developers
 - Export files
 - Develop interfaces
 - Import, store, and process data

Methods: Identifying Partners (Cont'd)

 Staff & a MU Stage 2 certified vendor provided a demonstration at the 2013 Missouri Dermatological Society Annual Meeting in St. Louis, MO

Methods: Software Issues

- Worked with other state and national groups/ organizations to:
 - Identify & assess software options that allow secure transfer of encrypted data via the Internet
 - MU's secure messaging software MoveIT (preferred)
 - Direct, PHIN-MS (acceptable)
 - Registry staff serve on national workgroups to develop/implement MU Stage 2 – Cancer Reporting guidelines
 - C/PO & Mapping Workgroups
 - Data elements
 - Formats
 - Triggers

Methods: Software Issues (Cont'd)

Added a specialty physician (urologist)
Trying two options:

Pros:	Trigger Event	Physician-driven
	Automated	Physician decides when to send
	More data	CCR gets critical data
		Easier to process at CCR



Methods: Registration of MU 2 Intent

- Collaborated with Missouri Department of Health and Senior Services staff to increase the number C/POs submitting EHR cancer data to MCR via DHSS's website for MU attestation and reporting
 - http://health.mo.gov/atoz/mophie/
 - http://mcr.umh.edu/mcr-meaningfuluse.php

Results: C/PO Participation (Cont'd)

- C/PO #1: Rural clinic completely electronic throughout:
 - Approached their EHR vendor (MediTech[™]) at HIMSS 2012
 - MediTech began working on changing reports to CDA formatted reports
 - Clinic developed implementation strategy of new cancer-reporting module
 - Received test data that was analyzed and feedback given to MediTech
 - Changes made to reports
 - EHR 2nd in country to be certified for MU Stage 2 Cancer Reporting by Office of National Coordinator (ONC) (Feb 2013)
 - MCR-ARC expected to receive live data Summer 2013
 - Revised date is Summer 2014

Results: Specialist C/PO Participation

C/PO #2: Urologist

- Joined project in 2012
- Received test data that was form-based EMR
 - Contacted EMR vendor (BuildYourEMR[™]) to adapt their reports for cancer-reporting to CDA formatted reports
- Received subsequent test data that was analyzed
 - BuildYourEMR changed some formatting issues
- EHR vendor 3rd in country to be ONC certified for MU Stage 2 Cancer Reporting (June 2013)

Results: Project Status

- Urologist Implementation has been completed
 - Live data anticipated soon
 - Analysis of data will begin immediately upon receipt of live data
 - Practice averages between 50-100 cases per year
 - Prostate cancers have never been received from a C/PO by MCR-ARC before

Results: CAH Participation

- Three CAHs
 - Selected EHR: 3
 - Implemented: 0

 Since none have implemented their EHRs, no preliminary findings

Results: Registration of MU 2 Intent

- DHSS is a centralized location for MU 2 reporting
 - Other than during the pilot, reports will be routed through DHSS
 - A similar process is used for ePath reports that MCR has been receiving for several years
- As of Jan 2014, 6 have registered their intent to participate in Stage 2 with DHSS

Challenge: Software

- Interoperability between C/PO and CCR software
- Convincing EHR vendors to change to CDA format before Stage 2 (1/1/14)
 - MU2 postponed a year during pilot and uncertainty of Cancer Reporting's inclusion
 - Convincing vendors to create a module for MU2 Cancer Reporting
- Convincing C/POs to choose cancer reporting as one of three options in MU Stage 2
 - Statutory, but no MU2 obligation

Challenge: C/PO Participation

- On-boarding additional C/POs
 - Targeted specialties
 - Need to determine #s
 - Other specialties that diagnose/treat cancer
 - Parts of state have few practitioners in targeted specialties
- Additional resources will be needed

Challenges: Staffing & Infrastructure

- Funding cuts
 - Staffing deficits (4 core positions)
 - Limits CCR's ability to implement EHR reporting by C/POs not in pilot
- Processing data and internal workflow
 - Storage
 - Consolidation of reports
- State HIE is under development
 - DHSS hopeful it would be up and going before 2015

Conclusions

- Identifying cost-effective ways for CCRs and non-hospital reporters to capture cases and report as mandated by law is challenging but rewarding
- Obstacles remain to be overcome but use of EHRs presents a viable solution
- Funding challenges remain
- Barrier: convincing C/POs to choose cancer reporting & convince EHR vendor to create the necessary module

Questions?

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