The Clinical Quality Framework Initiative to Harmonize Decision Support and Quality Measurement Standards

Defined Standards, Pilot Results, and Moving Beyond Quality Improvement

Panel Members

| Panelist | Title | Initiative Role |
|--|--|--|
| Julia Skapik, MD, MPH Julia.skapik@hhs.gov | Medical Officer Office of the National Coordinator for Health IT | Executive Sponsor |
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CQF Wiki: cqframework.info

Julia Skapik

CQF MOTIVATION AND GOALS

The Learning Healthcare System



 Build evidence out of practice

Set standards based on clinical goals, and evidence-based practice

Leverage EHR
to optimize
workflow and
support
clinical
decision
making

 Focus on "Making it Easy to do the Right Thing"

 Leverage analytics to extract actionable knowledge Measure the impact of the change through outcomes analysis and research

Develop reports to monitor the practice change

 Collaborate to foster knowledge translation

Meaningful Use is a Building Block



Use information to transform

Improve access to information

Data utilized to improve delivery and outcomes

Patient self management

Care coordination

Evidenced based medicine

Registries for

disease management **Privacy & security** protections

Connect to Public Health

Enhanced access and continuity

Data utilized to improve delivery and outcomes

Patient engaged, community resources

Patient centered care coordination

Team based care. case management

Registries to manage patient populations

Privacy & security protections

Connect to Public Health

Use technology to gather information

> **Basic EHR** functionality, structured data

Privacy & security protections

Connect to Public Health

Care coordination

Patient engaged

Connect to Public Health

Privacy & security protections

Structured data utilized for Quality **Improvement** Office of the National Coordinator for Health Information Technology

ACOs

PCMHs ACO
The Star and Swoo**Stage**g1h(MUealth IT, the Puttin**Stage**H2/l(MU)mposite logo, HealthIT.gov, the HealthIT.gov composition logo, HealthITBuzz composite logo are service marks or registered service marks of the U.S. Departme3:-ParthAimuman Services. Stage 3 hITBuzz, and the Stage 3 MU

2015 Edition Specific Health IT Goals



Improve Interoperability

Facilitate Data Access and Exchange

Ensure
Privacy and Security
Capabilities

Improve Patient Safety

Reduce Health Disparities

Improve the Reliability and Transparency of Certified Health IT

Use the ONC Health IT
Certification Program to
Support the Care Continuum

Support Stage 3 of the EHR Incentive Programs

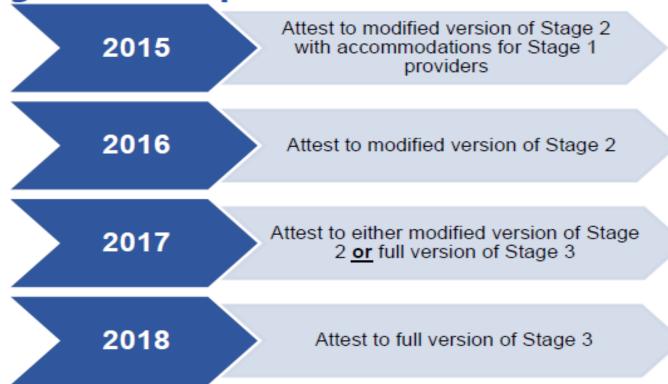
CMS MU Stage 3 Proposed Rule Reporting and Participation



Reporting Period

- Full calendar year reporting period beginning in 2017
- CQM reporting in coordination with quality reporting programs

Changes to Participation Timeline



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Improving Medicare Post-Acute Care Transformation (IMPACT) Act of 2014 Health IT.gov

- IMPACT Act added new section 1899(B) to Title XVIII of the Social Security Act (SSA)
- Post-Acute Care (PAC) providers must report:
 - Standardized assessment data
 - Data on quality measures
 - Data on resource use and other measures
- The data must be standardized and interoperable to allow for the:
 - Exchange of data using common standards and definitions
 - Facilitation of care coordination
 - Improvement of Medicare beneficiary outcomes
- PAC assessment instruments must be modified to:
 - Enable the submission of standardized data
 - Compare data across all applicable providers

IMPACT ACT: Quality Measure Domains



Requirements:

- Measures must be uniform/standardized across the 4 settings
- Measures will be risk adjusted, as determined appropriate by the Secretary

Domains:

- Functional status, cognitive function, and changes in function and cognitive function
- Skin integrity and changes in skin integrity
- Medication reconciliation
- Incidence of major falls
- Communicating the existence of and providing for the transfer of health information and care preferences

Appropriate Use Criteria for Advanced Diagnostic Imaging Services



- Section 218(b) of the PAMA amended Title XVIII of the Act, to establish a program to promote the use of appropriate use criteria (AUC) for advanced imaging services.
- The legislation requires in 2018 that every claim for advanced radiologic studies would include both:
 - Evidence that the user had utilized some form of approved clinical decision support that supported "appropriate use" of the advanced radiologic study
 - Evidence as to whether the user adhered to that advice or not

Appropriate Use Criteria for Advanced Diagnostic Imaging Services



- The goal of the PAMA legislation is to curb the ordering of unnecessary advanced radiologic studies, which make up billions of dollars of Medicare and Medicaid spending
- "Advanced radiologic studies" includes CT, MRI, fMRI, SPECT, PET and other nuclear studies but not traditional X-rays

Appropriate Use Criteria for Advanced Diagnostic Imaging Services



- CMS is currently establishing criteria for clinical "appropriate use"
- In 2016 they will establish how users can technically demonstrate the use of "appropriate use"
- In 2019 Congress mandated that CMS start to require preauthorization of radiologic studies for entities and individuals found to be regularly ordering studies deemed "inappropriate" according to the established "appropriate use" criteria



CMS quality reporting and performance programs

Hospital Quality

- •Medicare and Medicaid EHR Incentive Program
- PPS-Exempt Cancer Hospitals
- ·Inpatient Psychiatric Facilities
- ·Inpatient Quality Reporting
- •HAC reduction program
- •Readmission reduction program
- •Outpatient Quality Reporting
- •Ambulatory Surgical Centers

Physician Quality Reporting

- Medicare and Medicaid EHR Incentive Program
- PQRS
- eRx quality reporting

PAC and Other Setting Quality Reporting

- Inpatient Rehabilitation Facility
- Nursing Home Compare Measures
- LTCH Quality Reporting
- Hospice Quality Reporting
- Home Health Quality Reporting

Payment Model Reporting

- Medicare Shared Savings Program
- Hospital Value-based Purchasing
- Physician
 Feedback/
 Value-based
 Modifier*
- ESRD QIP

"Population" Quality Reporting

- Medicaid Adult Quality Reporting*
- CHIPRA Quality Reporting*
- Health
 Insurance
 Exchange
 Quality
 Reporting*
- Medicare Part
- Medicare Part D*

^{*} Denotes that the program did not meet the statutory inclusion criteria for pre-rulemaking, but was included to foster alignment of program measures.

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Mandated coordination across agencies and programs:

HHS Measurement Alignment



Measures

Unified

Outcome

Health IT.gov

current

EHR as primary reporting platform, with secondary reporting from registry, claims

MACRA: Medicare Access and CHIP Reauthorization Act of 2015



Beginning in 2019, all current Medicare payment, including incentive programs, will be combined into one Merit-Based Incentive Payment System (MIPS), replacing all Medicare reimbursement for eligible professionals.

The MIPS program will use four performance measures to determine reimbursement, which will begin in 2019:

- Quality;
- Resource use;
- Clinical practice improvement activities; and
- Meaningful use of certified EHR technology.

Privacy and security including HIPAA are also requirements and failure to adhere to required standards results in penalties

APMs & MIPS Paying for Performance



Alternative Payment Model (APM)

Clinicians who receive a substantial portion of their revenues (at least 25% of Medicare revenue in 2018-2019 but threshold will increase over time) from *qualifying* alternative payment mechanisms will not be subject to MIPS.

While the definition of a qualifying APM has yet to be determined, MACRA outlines criteria which includes but is not limited to:

Quality Measures Use of certified EHR technology

Risk-sharing

Merit-Based Incentive Payment System (MIPS)

Adjustments based on the **composite performance score** of each eligible physician or other health professional on a 0-100 point scale based on the following performance measures. All scores noted below are for the first MIPS year and are subject to adjustment. Additional positive adjustment available for exceptional performance.

| Quality (30% of MIPS score for first 2 years) | Clinical Practice Improvement Activities (15%) |
|---|---|
| Resource Use (10% 1st year) | Meaningful Use of certified HER (15%) |

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MIPS-Eligible Professionals (EP) Notable Dates



July 1, 2017

CMS must make available timely confidential feedback reports to each MIPS EP

Qualifying EPs 2019-20

- Physicians
- PAs
- Certified RN Anesthetists
- NPs
- Clinical Nurse Specialists
- Groups that include such professionals

2017

2018

2019

2020

2021

July 1, 2018

CMS must make available to each MIPS EP information about items and services furnished to the EP's patients by other providers and suppliers for which payment is made under Medicare

2021 & Onward

Secretary can add EPs to MIPS

MACRA: Medicare Access and CHIP Reauthorization Act of 2015



Failing to perform to the program minimums results in payment penalties:

- 2019- 4% maximum penalty
- 2020-5% maximum penalty
- 2021- 7% maximum penalty
- 2022-9% maximum penalty

Eligible professionals with higher performance scores receive an incentive up to three times the annual cap for negative payment adjustments

Nationwide interoperability is a requirement by December 31, 2018*

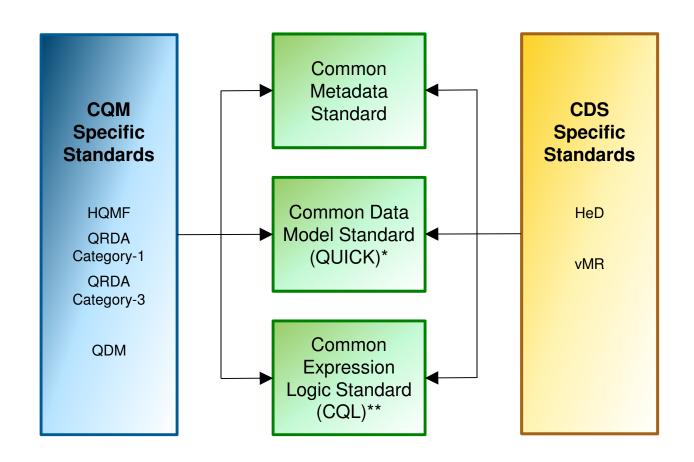
What's the connection to the Clinical Quality Framework?



- Current and Stage 3 MU rules require the use of CDS that supports at least 5 eCQMs
- Yet, the current eCQM constructs do not support CDS
- Future HHS programs will increasingly reference eCQMs and/or CDS
- Current standards are too complex and not computable enough
- New standards are emerging in the industry

Standards improvement and harmonization:

Clinical Quality Measurement and Clinical Decision Support Health IT.gov



^{*} Quality Improvement and Clinical Knowledge

^{**} Clinical Quality Language

The Current State vs the Future Vision



Current State

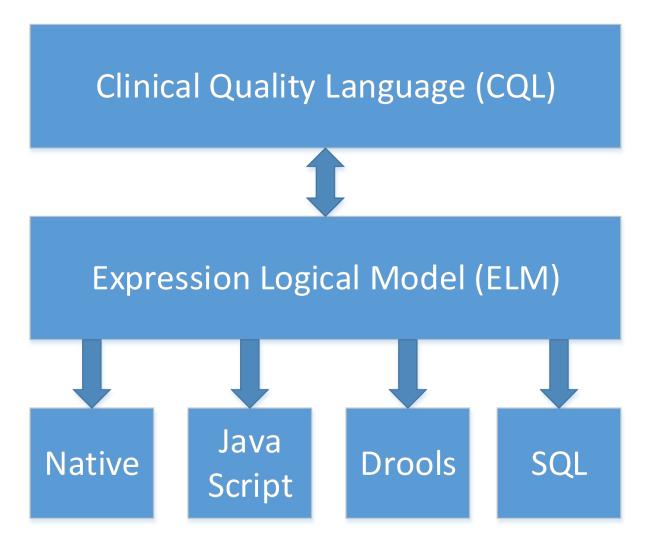
- CQMs and CDS are separate
- Each vendor develops their own CDS artifacts
- CQMs are focused on retrospective data
- CDS is an afterthought

Future Vision

- CDS drives care activities
- Performance is consistently improved through CDS
- CQM data capture is automatic
- CQMs are available with paired optional CDS artifacts

Future Standards Need to be Interoperable to Each Other to Allow Flexibility





Authors use CQL to produce libraries containing human-readable yet precise logic.

ELM XML documents contain machine-friendly rendering of the CQL logic. This is the intended mechanism for distribution of libraries.

Implementation
environments will either
directly execute the ELM, or
perform translation from
ELM to their target
environment language.

Moving Towards Next Generation Standards



- Staged approach to introduction of harmonized content starting with the Clinical Quality Language (CQL-based HQMF)
- Future HHS programs could allow optional FHIR reporting before requiring a transition
- Use of APIs and maps could facilitate consistent translation from one standard to another without loss of meaning

The Future Vision: Seamless Decision Support that Autogenerates Quality Data Health IT.gov

- Uses clearly defined, well-investigated data elements for capture at the point of care
- Allows patient-generated data to be integrated into measures and CDS
- Clinicians take no additional workflow steps to document quality metrics
- Decision support is educational, interactive, and facilitated by sophisticated data modeling



The Electronic Clinical Quality Improvement (eCQI) Resource Center



ecqi.healthit.gov

What is the eCQI Resource Center?

- The Resource Center is designed to act as a central hub for storing and collating resources surrounding the eCQMs and CDS standards, measures, tools, and guidance.
- It is cosponsored by CMS and ONC
- It will continue to add functionality and additional related content over time
 - We welcome your feedback!





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- Contact us at: onc.request@hhs.gov
- Visit the Health IT Dashboard: dashboard.healthit.gov
- Make speaker requests here:

http://www.healthit.gov/requestspeaker

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http://www.youtube.com/user/HHSONC

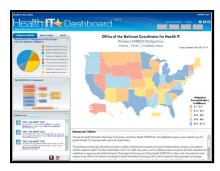
Linked in Health IT and Electronic Health Records

http://www.scribd.com/HealthIT/

flickr http://www.flickr.com/photos/healthit

Providers & Professional

| Providers & Professional | Professional | Policy Researchers |





Health IT Buzz Blog

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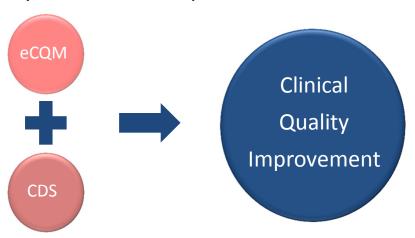
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Marc Hadley

CQF STANDARDS

Background

- Clinical Decision Support (CDS) and electronic Clinical Quality Measurement (eCQM) are closely related, share many common requirements, and both support improving health care quality.
 - CDS guides a clinician to follow a standard plan of care
 - eCQM measures adherence to a standard plan of care
- Shared needs:
 - Define patient cohorts (sub-populations)
 - Standard ways to reference patient data in EHR



The Challenge

- Current eCQM and CDS standards
 - were not developed together
 - use different approaches to patient data
 - use different approaches to expression logic

| | References to Patient Data | Expression Logic | Exchangeable Artifacts |
|------|------------------------------|------------------------------|------------------------------|
| CDS | Virtual Medical Record (VMR) | CDS Knowledge Artifact (HeD) | CDS Knowledge Artifact (HeD) |
| eCQM | Quality Data Model (QDM) | Quality Data Model (QDM) | QRDA I & III, HQMF |

EHR vendors and homegrown systems must

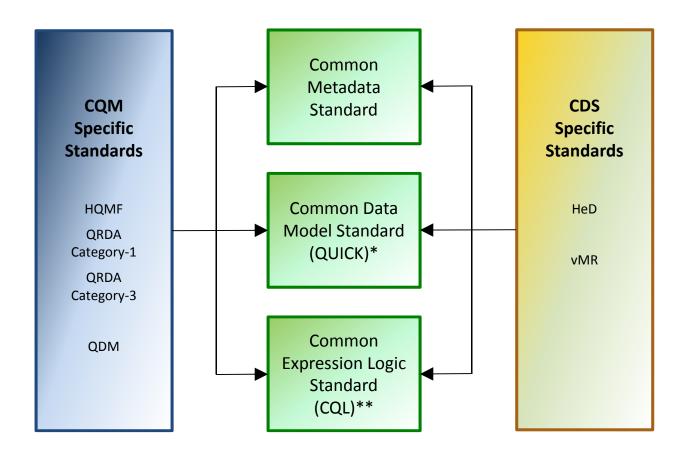
- Map their data to two different data model standards
- Implement computation of two different logic standards
- Interpret and implement text "guidance"

eCQM and CDS rule authors cannot

- Share or reuse logic between measures and rules
- Ensure consistency between matching measures and rules
- Adequately express all of their requirements

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The Goal: Shared Standards



^{*} Quality Improvement and Clinical Knowledge

^{**} Clinical Quality Language

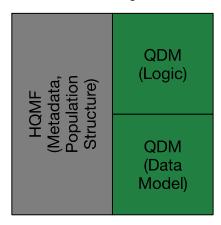
Standards Harmonization Impact

- Improves efficiency and reduces cost
 - eCQM / CDS system implementation
 - eCQM / CDS rule authoring and maintenance
- Improves consistency and accuracy
 - Shared logic between measures and rules
 - Reduce or eliminate need for "guidance"
- Improves quality of standards
 - Leverage past lessons learned from eCQM & CDS
 - Community effort from larger, more diverse community
- Promotes integration of CQM and CDS domains

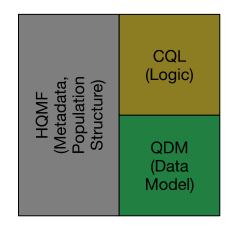
Bottom Line: Improves the Quality of Care Patients Receive

Evolving eCQM Standards

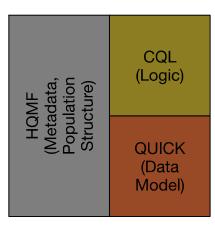
Today



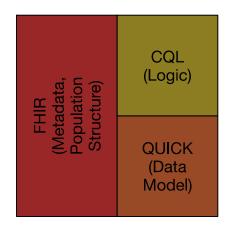
Near term solution



Rejected



Long term solution



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A shared expression language for eCQM and CDS

CLINICAL QUALITY LANGUAGE

What is the Clinical Quality Language?

"The Clinical Quality Language Specification defines a representation for the expression of clinical knowledge that can be used within both the Clinical Decision Support (CDS) and Clinical Quality Measurement (CQM) domains."

HL7 Standard: Clinical Quality Language Specification, DSTU Release 1

CQL Key Points

- The CQL specification defines two components:
 - Clinical Quality Language: Author-friendly domain specific language
 - Expression Logical Model: Computable XML
- CQL leverages best practices and lessons learned from:
 - Quality Data Model: Focus on ease of authoring
 - Health eDecisions: Focus on modularity and computability
 - eCQM & CDS Communities: HL7 Work Groups and S&I Framework
- CQL is designed to work with any data model
- CQL is much more expressive and robust than QDM logic
- CQL is a Health Level 7 (HL7) Draft Standard for Trial Use (DSTU)

Example: CMS 123 – Diabetes: Foot Exam

Initial Population

Patients 18-75 years of age with diabetes with a visit during the measurement period

```
define InInitialPopulation:
```

```
AgeInYearsAt (start of MeasurementPeriod) >= 18

and AgeInYearsAt (start of MeasurementPeriod) < 75

and exists (["Diagnosis": "Diabetes"] D where D.period overlaps MeasurementPeriod)

and exists (ValidEncounters E where E.period during MeasurementPeriod)
```

define ValidEncounters:

```
["Encounter, Performed": "Office Visit"]
union ["Encounter, Performed": "Face-to-Face Interaction"]
union ["Encounter, Performed": "Preventive Care Services Established Office Visit"]
union ["Encounter, Performed": "Preventive Care Services Initial Office Visit"]
union ["Encounter, Performed": "Home Healthcare Services"]
```

Denominator

Equals Initial Population

define InDenominator:

InInitialPopulation

Example: CMS 123 – Diabetes: Foot Exam

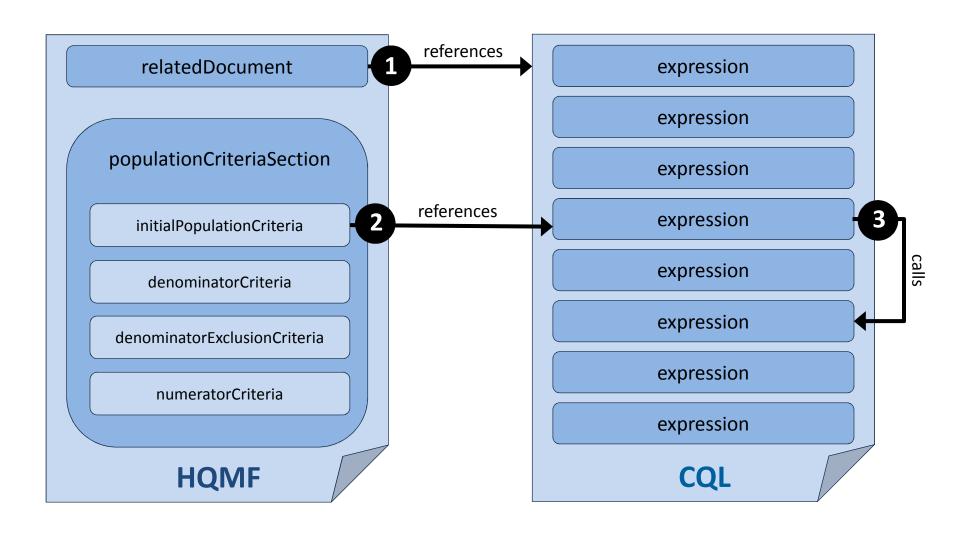
Numerator

Patients who received visual, pulse and sensory foot examinations during the measurement period

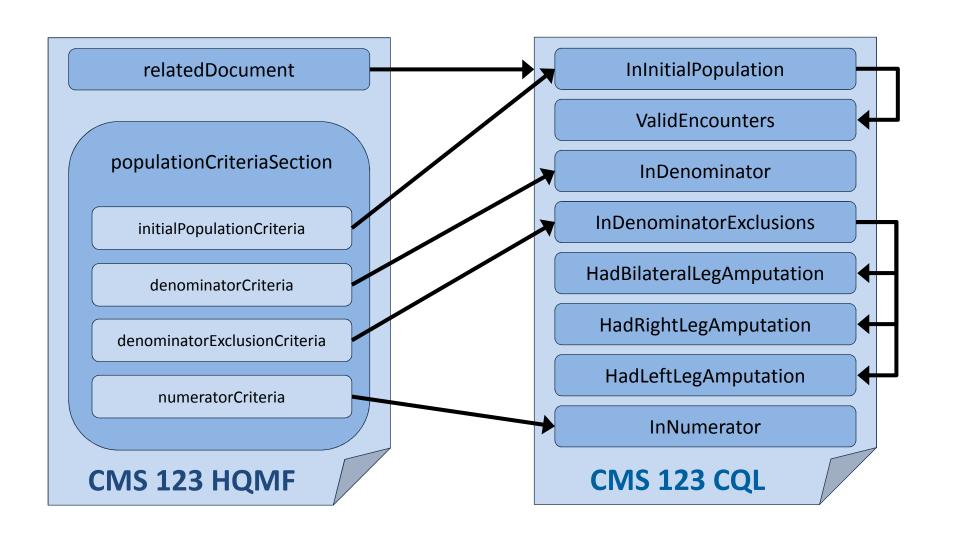
define InNumerator:

```
exists (["Procedure, Performed": "Visual Exam of Foot"] P
  where P.period during MeasurementPeriod)
and exists(["Procedure, Performed": "Sensory Exam of Foot"] P
  where P.period during MeasurementPeriod)
and exists (["Procedure, Performed": "Pulse Exam of Foot"] P
  where P.period during MeasurementPeriod)
```

CQL-based HQMF



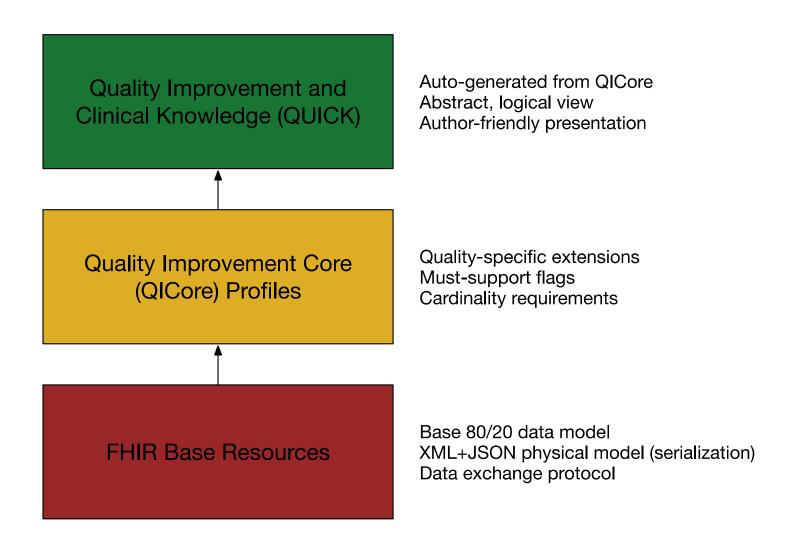
CQL-based HQMF for CMS 123

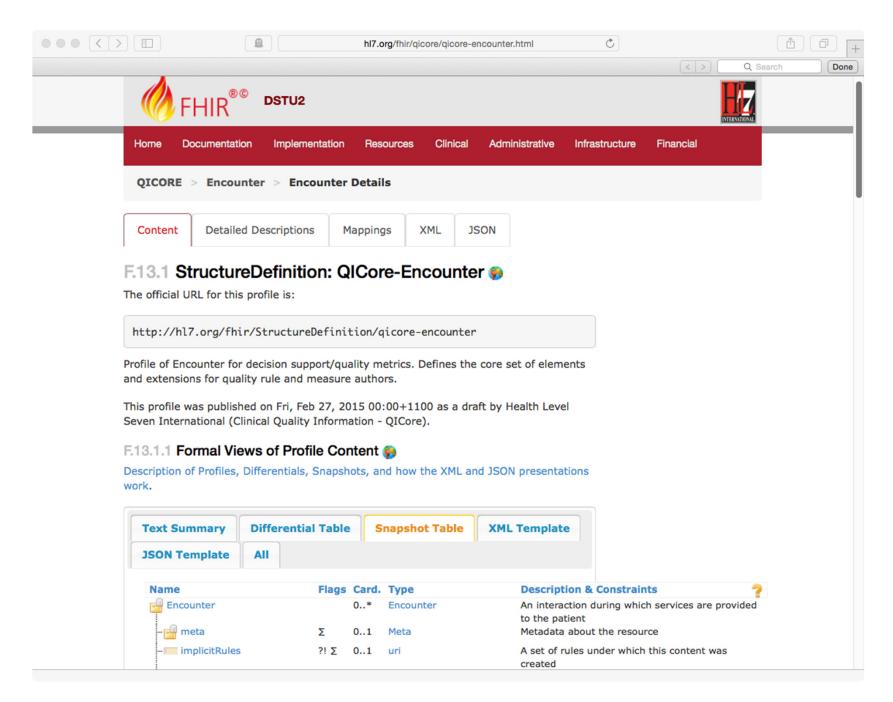


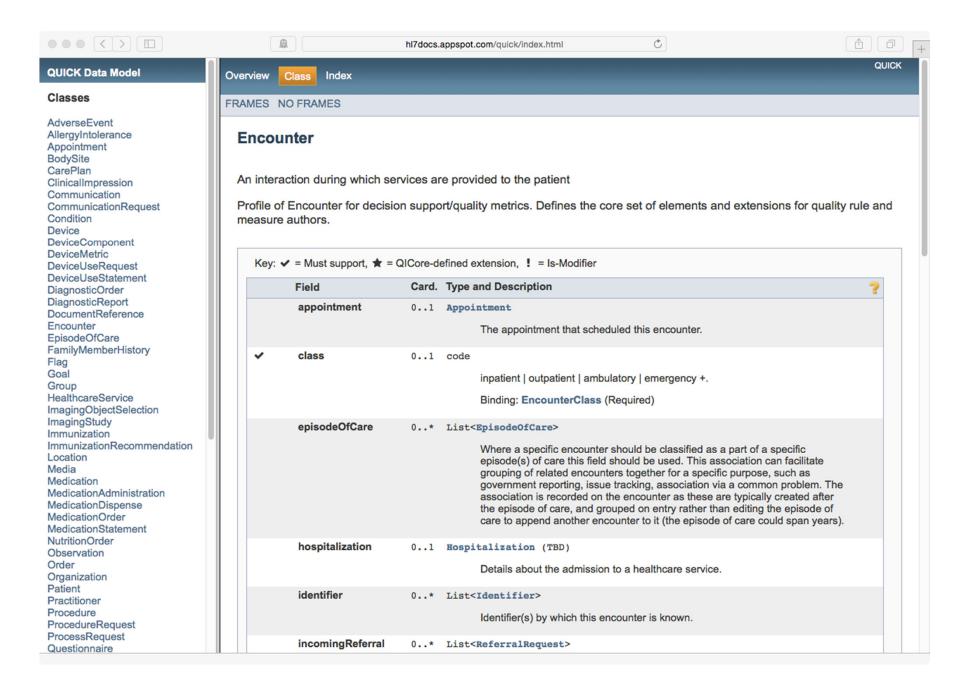
A shared data model for eCQM and CDS

QUALITY IMPROVEMENT AND CLINICAL KNOWLEDGE (QUICK)

OIIICK Derivation

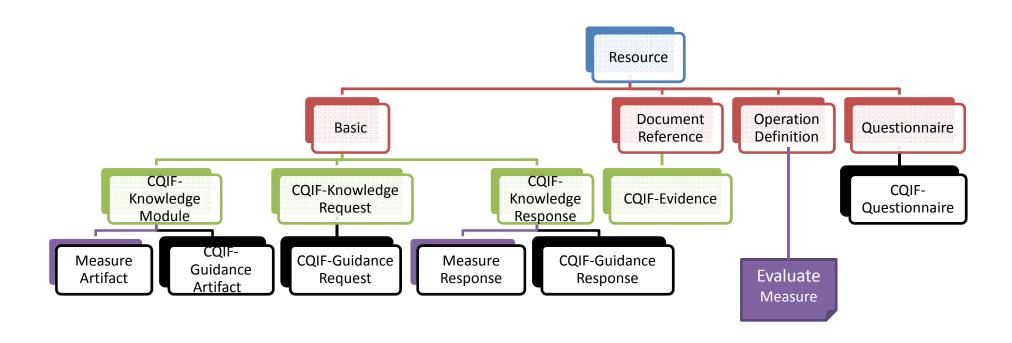






FHIR-BASED ECQMS AND CDS

eCQM and CDS Profiles



Standards Status

- CQL
 - Draft standard at HL7
- CQL-based HQMF IG
 - Draft standard at HL7
- CQL-based Health eDecisions Knowledge Artifact HeD KA R1.3
 - Draft standard at HL7
- QICore
 - Draft standards, part of FHIR DSTU R2
- QUICK
 - To be published in a DSTU update of QICore
- CDS FHIR Profiles and eCQM FHIR Profiles
 - Balloted "for comment" in Sept HL7 ballot cycle
 - Planning for draft standard ballot in May HL7 ballot cycle

Resources

- CQL Specification
 - http://www.hl7.org/implement/standards/product_brief.cfm?product_id=400
- CQL-based HQMF Implementation Guide
 - http://www.hl7.org/implement/standards/product_brief.cfm?product_id=405
- CDS/eCQM Harmonization on the eCQI Resource Center
 - https://ecqi.healthit.gov/cdsecqm-harmonization
- S&I Clinical Quality Framework Initiative
 - http://wiki.siframework.org/Clinical+Quality+Framework+Initiative
- HL7 CQI Work Group
 - http://www.hl7.org/special/committees/CQI/index.cfm
- HL7 CDS Work Group
 - https://www.hl7.org/Special/committees/dss/index.cfm

Kensaku Kawamoto

PILOTS AND PARTNERSHIPS

Disclosures

- I am, or have been in the past, a consultant on clinical decision support to the following entities:
 - Office of the National Coordinator for Health IT (ONC)
 - Mayo Clinic
 - Partners HealthCare
 - McKesson InterQual
 - RAND Corporation
 - ARUP Laboratories

- ESAC, Inc.
- JBS International, Inc.
- Clinica Software, Inc.
- Religent, Inc.
- Inflexxion, Inc.
- Intelligent Automation, Inc.

| Pilots | Point of | Liaison SME |
|--|---------------------------|--------------|
| | Contact | |
| Breast Cancer Decision Support and | Chad Armstrong | Claude Nanjo |
| Clinical Quality Measurement (CQM) | | |
| Cardiology Appropriateness of Use | Rachel Davis | Chris Moesel |
| Chlamydia Screening | Johanna Goderre- Jones | Bryn Rhodes |
| Immunization Decision Support Services (DSS) | Daryl Chertcoff | Claude Nanjo |
| Phenotype Execution and Modeling | Will Thompson | Bryn Rhodes/ |
| Architecture | | Chris Moesel |
| Portable CDS Knowledge Artifacts | Julie Scherer | Claude Nanjo |
| Radiology Appropriateness of Use | Tom Conti | Bryn Rhodes |
| Others in Planning | TBD | TBD |
| (e.g., Opioid Management) | | |

Survey Results



100%

percentage of pilots that met their goal(s)



10+

artifacts and tools created as part of the pilots



2700+

hours spent by pilot teams in 4 months of pilot work



12:30:47

MORE TIME

resources that could have made pilots more successful



ALL

pilots felt they received the support they needed



8.4

average rating for overall CQF pilot experience

The project team has provided us with valuable guidance on the current state of the CQF standards, and how they will evolve moving forward. This is the kind of support we were looking for in joining as a pilot project.

...having one-on-one sessions with our pilot advisor Bryn really helped solidify/speed up our pilot design and progress...



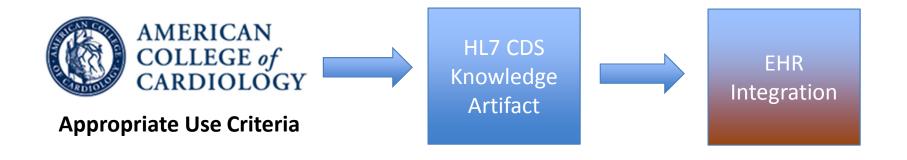
We see great opportunity and need for continued development of the standards...

Cardiology Appropriateness of Use

<u>Goal</u>: Provide point-of-order guidance on appropriate diagnosis and risk assessment of stable ischemic heart disease based on American College of Cardiology (ACC) Appropriate Use Criteria

| Team Member | Role |
|--|------------------|
| Rachel Davis, MS | ACC SME and |
| Director of Health Information Technology, ACC | Point of Contact |
| James Tcheng, MD | ACC SME and |
| Professor, Duke Univ.; Chair, ACC Digital Steering Committee | Pilot Lead |
| Dino Damalas, MBA | ACC SME |
| CIO, ACC | |
| Joseph Allen | ACC SME |
| Director of Translating Research into Practice, ACC | |
| Ganesan Srinivasan, MBA | ACC SME |
| Director of Registry & BI Product Development, ACC | |
| Christopher Moesel | Liaison SME |
| Principal Computer Science Engineer, MITRE | |

Cardiology Appropriateness of Use



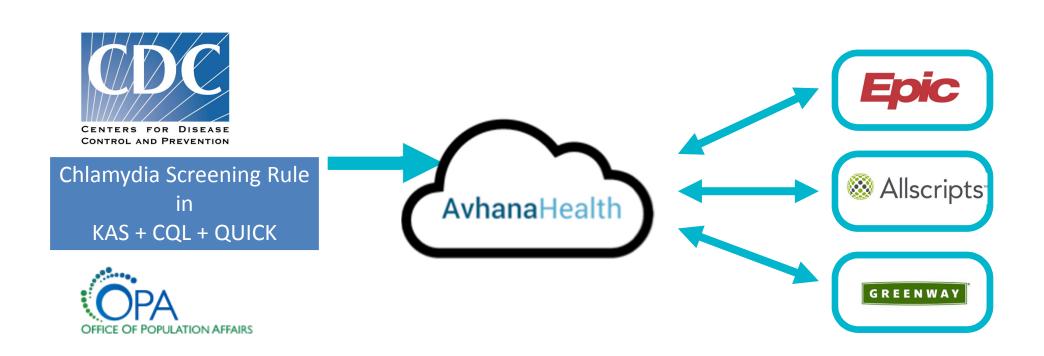
EHR=Electronic Health Record CQF Wiki: cqframework.info

Chlamydia Screening

<u>Goal</u>: Support CDS and eCQM for screening, treatment, and followup of *chlamydia trachomatis* infection in community settings

| Team Member | Role |
|---|------------------|
| Johanna Goderre Jones, MPH | Pilot Lead and |
| Senior Health Informatics Advisor, HHS Office of Population Affairs | Point of Contact |
| Noah Weiner | Technical Lead |
| CEO and Co-Founder, Avhana Health | |
| Lorrie Gavin, PhD, MPH | CDC Advisor |
| Health Scientist, CDC Division of Reproductive Health | |
| Victor Lee, MD | SME |
| VP, Clinical Informatics, Zynx Health | |
| Bryn Rhodes | Liaison SME |
| Owner, Database Consulting Group LLC | |

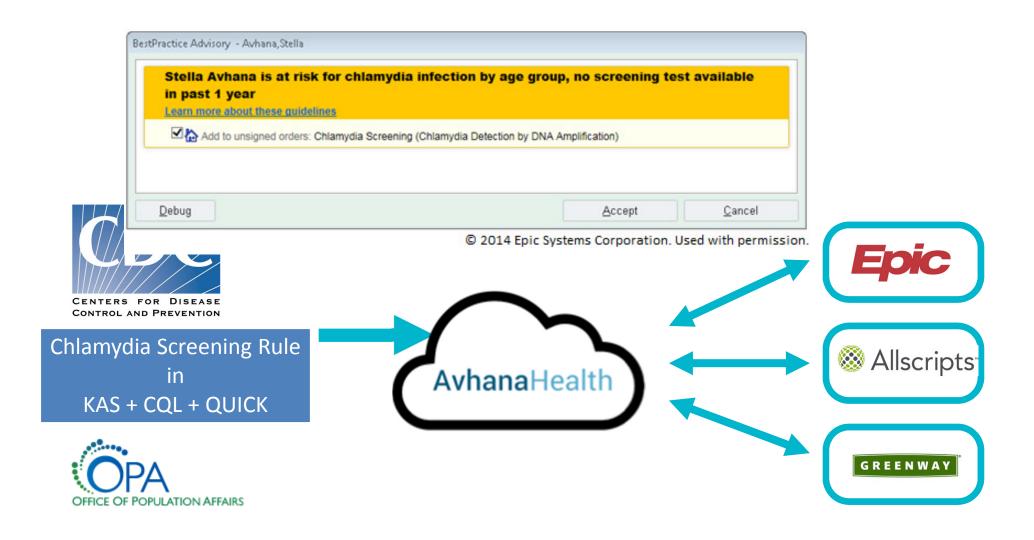
Chlamydia Screening



Chlamydia Screening



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Immunization Decision Support Service

<u>Goal</u>: Validate how CQF standards may support and evolve Clinical Decision Support for Immunizations (CDSi)

| Team Member | Role |
|--|-------------------|
| Daryl Chertcoff | Pilot Lead |
| Project Manager, HLN Consulting, LLC | |
| Eric Larson | Pilot Participant |
| American Immunization Registry Association; CDC Contractor | |
| Nathan Bunker | Pilot Participant |
| Senior Technical Project Manager, American Immunization | |
| Registry Association | |
| Mario Hyland | Pilot Participant |
| Senior Vice President, AEGIS.net, Inc. | |
| Richard Ettema | Pilot Participant |
| Lead Consulting, AEGIS.net, Inc. | |
| Claude Nanjo, MPH, MAAS | Liaison SME |
| Chief Scientist and Senior Software Architect, Cognitive Medical | |
| Systems, Inc. | |

Immunization Decision Support Service

Sample Immunization Forecasting User Interface



Immunization Decision Support Service

Validate CQF standards for Immunization Forecasting

Pilot accomplishments to date:

- Participated in 2 HL7 Connectathons
 - May 2014: Demonstrated immunization forecasting using CDSi logic, FHIR DSTU1 interface, and 1 immunization forecaster, with AEGIS WildFHIR serving as integration layer
 - January 2015: Demonstrated immunization forecasting using CDSi logic, FHIR DSTU2 interface, and 2 immunization forecasters
- July 2015: Shared lessons learned with CQF as feedback for development of FHIR-based update to DSS standard
- In discussions for participation in next HL7 Connectathon

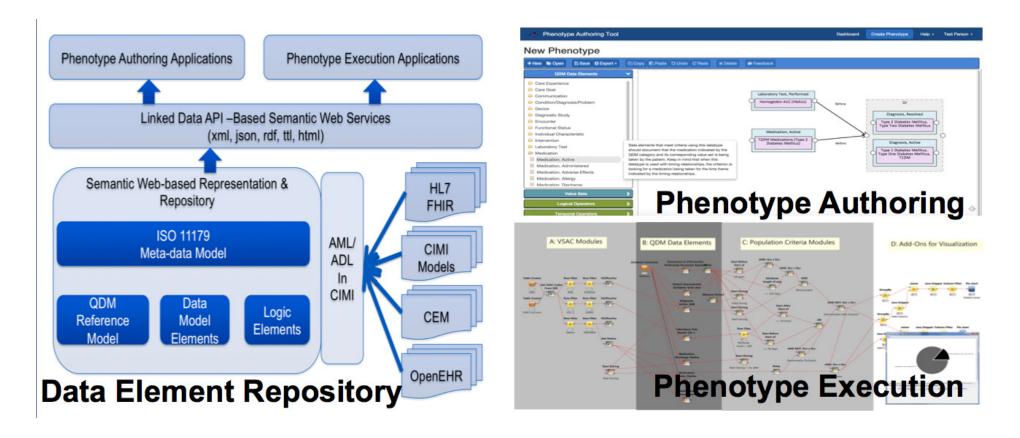
Phenotype Execution and Modeling Architecture (projectphema.org; R01 GM105688)

<u>Goal</u>: Take HQMF/QDM-based Phenotype Authoring, Execution, and Knowledge Management Platform and enable support for CQF standards

| Team Member | Role |
|---|------------------|
| William Thompson, PhD | Pilot Lead and |
| Research Scientist and Adjunct Assistant Professor | Point of Contact |
| Northwestern University & NorthShore University HealthSystem | |
| Jyotishman Pathak, PhD | Pilot Co-Lead |
| Professor and Chief | |
| Division of Health Informatics, Weill Cornell Medical College | |
| Bryn Rhodes | Liaison SME |
| Owner, Database Consulting Group LLC | |
| Christopher Moesel | Liaison SME |
| Principal Computer Science Engineer, MITRE | |

Phenotype Execution and Modeling Architecture

(projectphema.org; R01 GM105688)



November 16th; Session S32: PheMA Demonstration November 17th; Session S68: PheMA Paper Presentation November 18th; Session S92: Phenotyping Panel

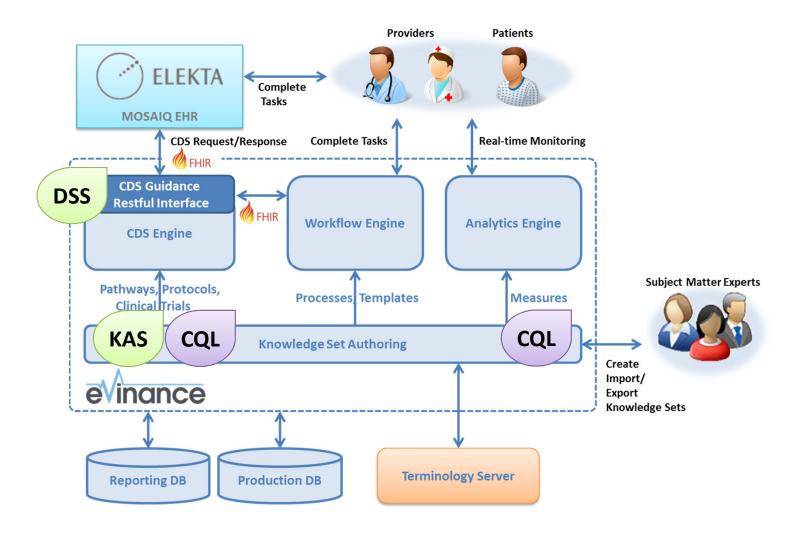
Breast Cancer Decision Support and CQM

Goals:

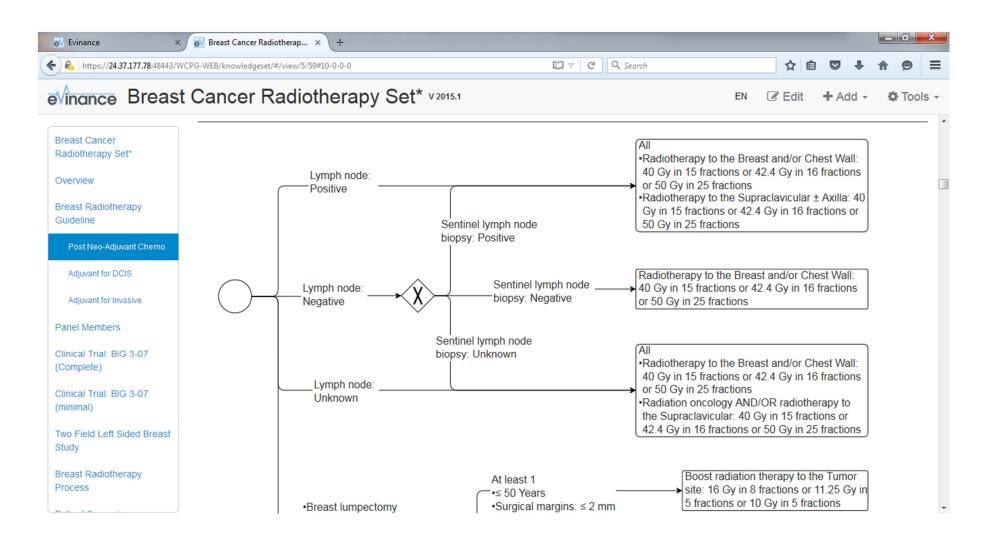
- 1. Demonstrate that FHIR can be used in oncology to send patient data to a Decision Support Service, which in turn can recommend appropriate treatment plan(s) and clinical trial(s)
- 2. Demonstrate that oncology measures can be defined using CQL and KAS

| Team Member | Role |
|--|-------------|
| Chad Armstrong, MBA | Pilot Lead |
| CEO, Evinance | |
| Claude Nanjo, MPH, MAAS | Liaison SME |
| Chief Scientist and Senior Software Architect, Cognitive Medical | |
| Systems, Inc. | |
| Bryn Rhodes | Liaison SME |
| Owner, Database Consulting Group LLC | |

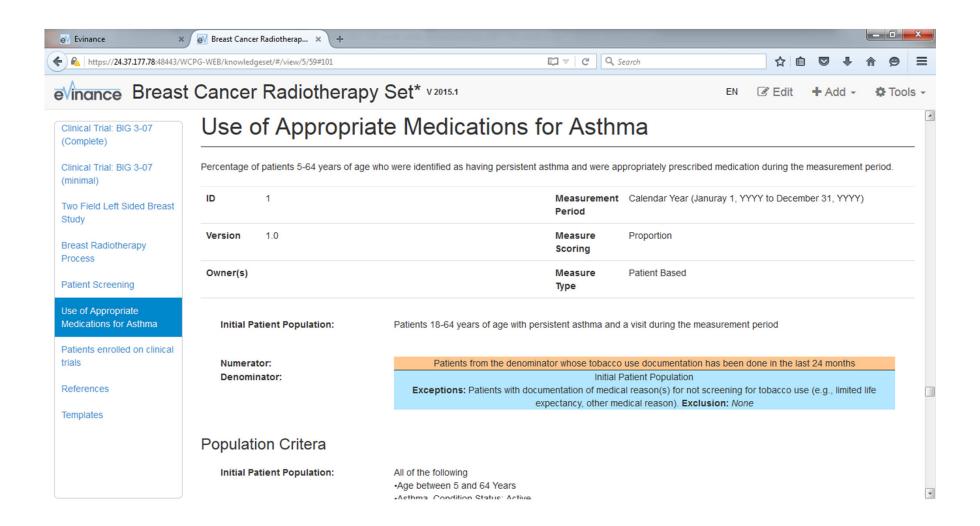
Technology Overview



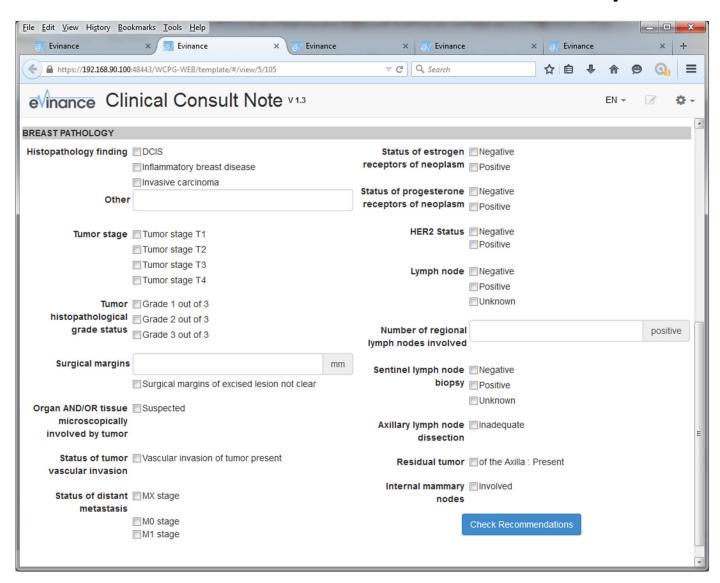
Guideline Authoring



Measure Authoring



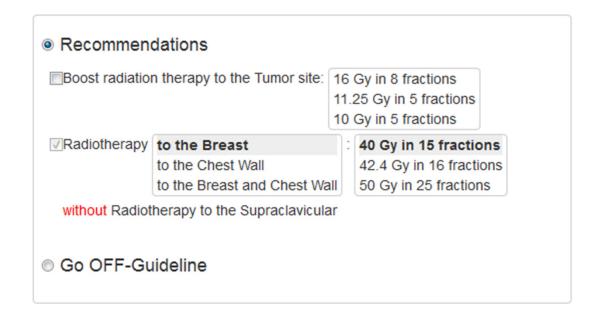
Structured Documentation Template



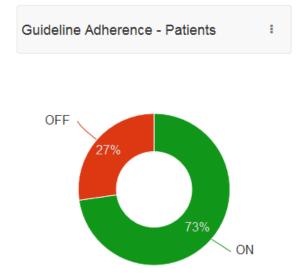
Care Recommendations - Prompts

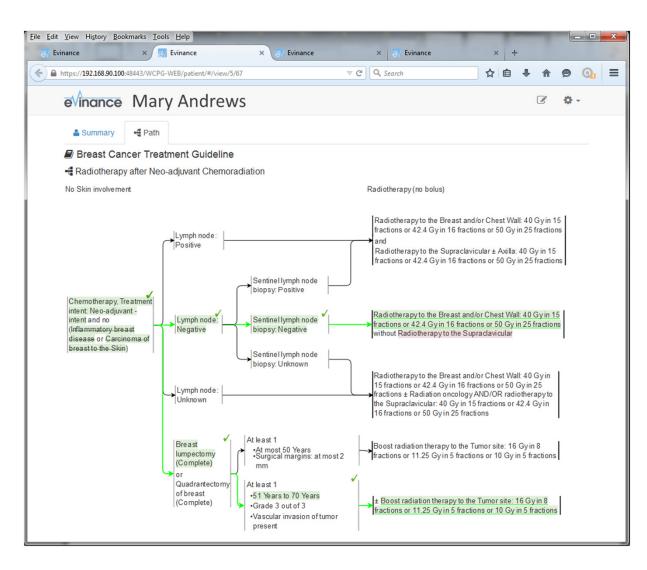


Care Recommendations



Guideline Adherence Dashboard







evinance community

evinance.com/community

Radiology Decision Support

<u>Goal</u>: Evaluate ability of CQF standards to support point-of-care, service-based, EHR-integrated CDS on radiology appropriateness of use in support of PAMA Section 218b provisions

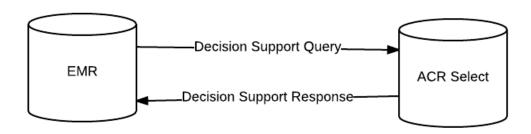
| Team Member | Role |
|---|------------------|
| Tom Conti | Pilot Co-Lead & |
| SVP Technical Strategy, National Decision Support Company | Point of Contact |
| Bob Cooke | Pilot Co-Lead |
| VP Marketing, National Decision Support Company | |
| James Doyle | SME |
| R&D Product Lead, Epic | |
| Erik Abels, MBA | SME |
| Clinical & Diagnostic Imaging Strategist, Cerner | |
| Keith Dreyer, DO, PhD | SME |
| Vice Chairman and Associate Professor of Radiology, MGH/Harvard | |
| Bryn Rhodes | Liaison SME |
| Owner, Database Consulting Group LLC | |

Background

- Leverage a widely deployed, commercial implementation of Radiology CDS
 - Currently process over 400,000 radiology decision support sessions per month
- Market implementations in place with Epic, Cerner, Meditech, McKesson, etc.
 - Leverage native EHR user interface for capture of structured reason for exam and display of feedback
 - All feedback is actionable (e.g., user can change/alter to more appropriate action)
- Embed transaction data within EHR including a unique Decision Support Number (DSN)

Pilot Architectural Diagram

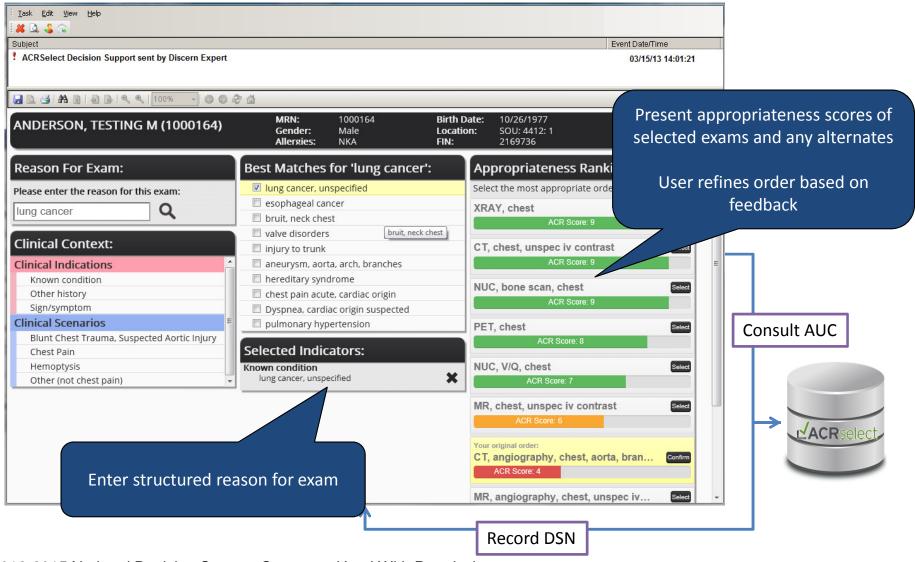
Current Integration



Pilot Integration



EHR Access to CDS Knowledge at Point of Care



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CQF Wiki: cqframework.info

Pilot Status

- Leveraged FHIR messaging and data standards
 - IHE Patient Care Coordination GAO (Guideline Appropriate Ordering)
 - http://ihe.net/uploadedFiles/Documents/PCC/IHE PCC
 Suppl GAO Rev1.0 PC 2015-06-01.pdf
 - GAO being harmonized with CQIF standard
- Translated existing Epic integration to use this new FHIR-based integration

CQF Pilots

Portable CDS Knowledge Artifacts

<u>Goal</u>: Demonstrate the authoring, translation, and accurate deployment of portable CDS knowledge artifacts represented in the CQF standard

| Team Member | Role |
|--|-------------|
| Julie Scherer, PhD | Pilot Lead |
| Chief Data Scientist, Motive Medical Intelligence | |
| Hanh Le, MD | SME |
| Director of Implementation, Motive Medical Intelligence | |
| Aziz Boxwala, MD, PhD | Liaison SME |
| CEO, Meliorix Inc. | |
| Claude Nanjo, MPH, MAAS | Liaison SME |
| Chief Scientist and Senior Software Architect, Cognitive Medical | |
| Systems, Inc. | |
| Rocky Reston, MD, PhD | SME |
| Chief Medical Informatics Officer, Cognitive Medical Systems, Inc. | |

CQF Wiki: cqframework.info

Pilot Objectives and Scope

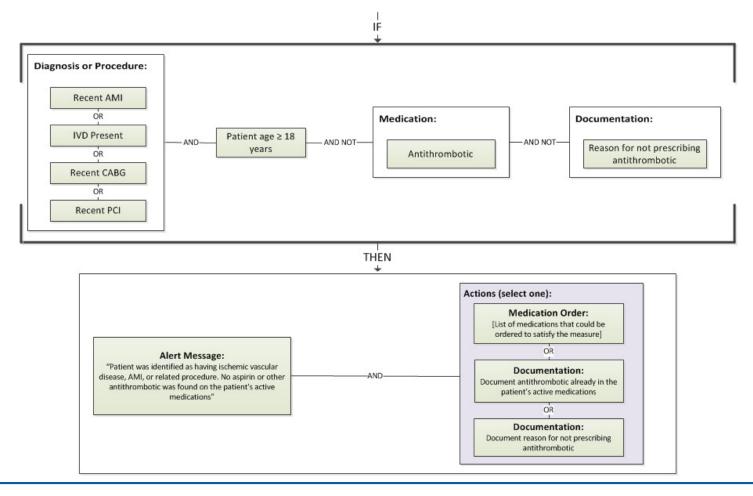
Pilot objectives

- Validate expressivity of the QUICK, QICore, and FHIR data models
- Validate expressivity of the CQL expression language
- Assess feasibility of artifact translation and execution
- Evaluate retention of artifact semantics and presentation behavior during translation and deployment

Implementation scope

- ECA rule based on eCQM 164, Ischemic Vascular Disease (IVD): Use of Aspirin or another Antithrombotic
- Documentation template of a suicide risk assessment
- Suicide risk assessment scoring logic
- Order sets for management and treatment of suicide risk

ECA Rule: Artifact Specification

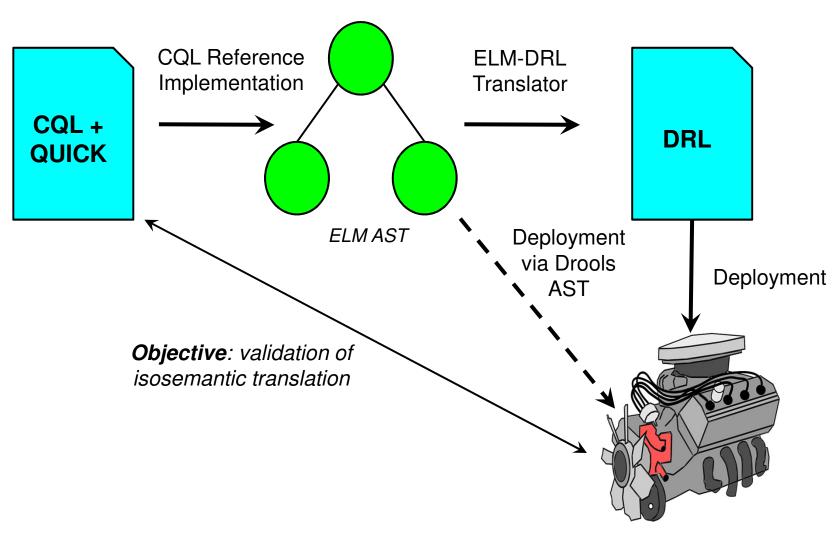


Diagrammatic representation of an ECA artifact for eCQM 164, Ischemic Vascular Disease (IVD): Use of Aspirin or another Antithrombotic

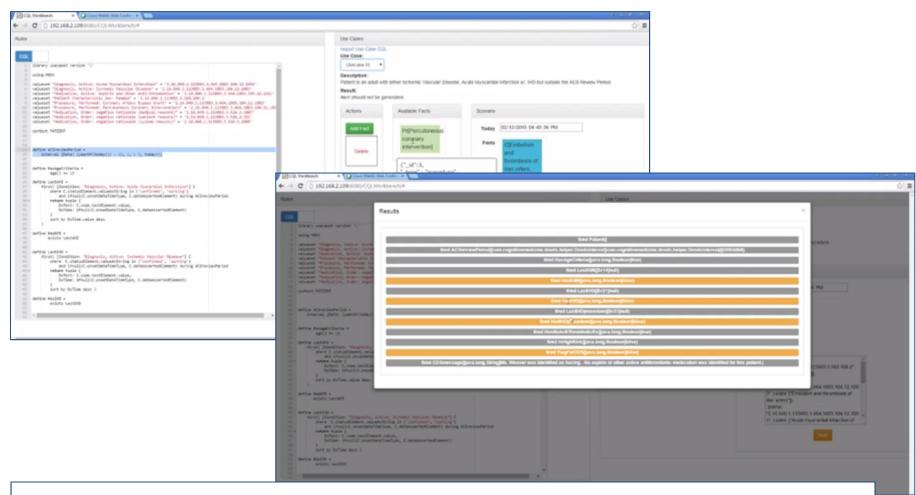
ECA Rule: CQL Artifact Excerpt

```
define LastAMI =
  first([Condition: "Diagnosis, Active: Acute Myocardial Infarction"] C
    where C.statusElement.valueAsString in {'confirmed', 'working'} and ifnull(C.onsetDateTimeType,
            C.dateAssertedElement) during ACSreviewPeriod
    return tuple { DxText: C.code.textElement.value, DxTime: ifnull(C.onsetDateTimeType,
            C.dateAssertedElement)
    sort by DxTime.value desc
define HadAMI = exists LastAMI
define LastIHDprocedure =
  first( ( [Procedure: "Procedure, Performed: Percutaneous Coronary Interventions"]
       union
       [Procedure: "Procedure, Performed: Coronary Artery Bypass Graft"] ) P
       where startOf(P."date") during ACSreviewPeriod
       return tuple { PxText: P.type.text,
               PxTime: startOf(P."date") }
         sort by PxTime.value desc )
define HadIHDprocedure = exists LastIHDprocedure
```

ECA Rule: Translation and Deployment

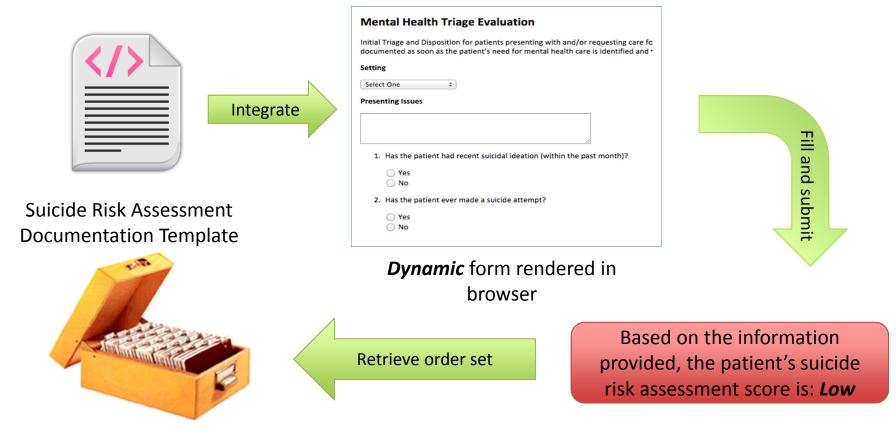


ECA Rule: Demonstration



Demonstration video available at https://vimeo.com/121041900 and begins at 36 minutes and 30 seconds

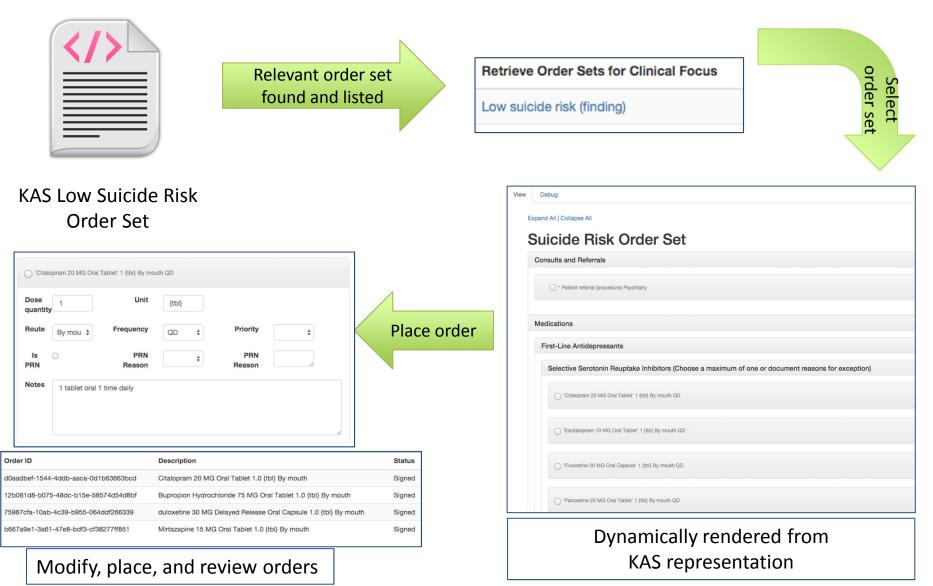
Documentation Template: Scoring and Order Set Identification



Suicide risk assessment score used to retrieve relevant KAS order set

ISON, executed upon form submission using Javascript

Order Sets: Retrieval, Presentation, and Submission



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CQF Wiki: cqframework.info

CQF Pilots

Key Insights from Pilots

- CQF standards can be leveraged to support clinically meaningful, real-world quality improvement use cases
- Detailed clinical models and associated value sets will need to be defined to ensure interoperability
- Specific vs. general interoperability specifications (e.g., GAO vs. CQIF) have tradeoffs and need to be harmonized
- Ease of implementation and alignment with industry trends (e.g., FHIR) will be critical to adoption
- Coordination and harmonization with other relevant initiatives will also be critical to adoption

CQF Pilots

Further Information

- Pilot lead contacts
 - cqframework.info
 - kensaku.kawamoto@utah.edu
- Pilot Showcase
 - August 27th, 2015
 - 200+ attendees
 - Materials and Video available at <u>http://wiki.siframework.org/Clinical+Quality+Framework+Past</u> <u>+Meetings</u>

Partnerships with Related Initiatives

- Goals:
 - Broaden the stakeholder community
 - Reduce implementer burden
- ONC Structured Data Capture (SDC) initiative
 - Collaborating to ensure alignment between CQIF documentation templates and SDC Questionnaires
- ONC Data Access Framework (DAF) initiative
 - QICore based on DAF FHIR profiles
 - Exploring options for closer alignment
- Clinical Information Modeling Initiative (CIMI) and Health Services Platform Consortium (HSPC)
 - Exploring whether CIMI/HSPC data models and FHIR profiles can be leveraged to meet CQF data needs

Tom Oniki

HSPC AND CIMI COLLABORATION

The Motivation for HSPC and CIMI

The complexity of modern medicine exceeds the inherent limitations of the unaided human mind.

- David M. Eddy, MD, Ph.D.

Intermountain and other providers can only provide the highest quality, lowest cost health care with the use of advanced clinical decision support systems integrated into frontline clinical workflow

But we can't keep up!

- We have ~150 decision support rules or modules
- We have picked the low hanging fruit
- There is a need to have 5,000 decision support rules or modules
- There is no path from 150 to get to 5,000 unless we fundamentally change the ecosystem

Strategic Goal

- We need a way to share what we create, benefit from what others create
- This includes:
 - data
 - applications
 - reports
 - alerts
 - protocols
 - decision support modules
- And effective sharing requires "plug-n-play" interoperability

What Is Needed to Create a New Paradigm?

- Standard set of detailed clinical data models coupled with standard coded terminology
- Standard APIs for healthcare related services
- Open sharing of models, coded terms, and APIs
- Sharing of decision logic and applications



Clinical Information Modeling Initiative

http://www.opencimi.org/

CIMI

- A community of interest that is producing detailed clinical models to enable interoperability of health care information systems
- Became an HL7 working group Oct 2015
 - interim co-chairs: Stan Huff, Intermountain; Virginia Riehl; Linda Bird, IHTSDO; Harold Solbrig, Mayo
- CIMI models are free for use for all purposes

Clinical modeling activities

- Netherlands/ISO Standard
- ISO EN 13606
- UK NHS and LRA
- Singapore
- Sweden
- Australia
- openEHR Foundation
- Canada
- US Veterans Administration
- US Department of Defense
- Intermountain Healthcare
- Mayo Clinic
- MLHIM

- SemanticHealthNet
- HI7
 - Version 3 RIM, message templates
 - TermInfo
 - CDA plus Templates
 - Detailed Clinical Models
 - greenCDA
- Tolven
- NIH/NCI (Common Data Elements, CaBIG)
- CDISC SHARE
- Korea CCM
- Brazil
- Others . . .

SystolicBPObs

code: LOINC code for "systolic blood pressure"

data: numeric + "mmHg"

qualifiers

BodyLocation (0..1)

data: <set of SNOMED codes for BP body locations>

PatientPosition (0..1)

data: <set of SNOMED codes for BP body positions>

SuspectedLungCancer

data: SNOMED code for "cancer"

qualifiers

BodyLocation (1)

data: SNOMED code for "lung"

Certainty (1)

data: SNOMED code for "suspected"

Stage (0..1)

What Is Needed to Create a New Paradigm?

- **Standard** set of detailed clinical data models coupled with...
- Standard coded terminology
- Standard APIs for healthcare related services
- Open sharing of models, coded terms, and APIs
- Sharing of decision logic and applications

SuspectedLungCancer

data: SNOMED code for "cancer"

qualifiers

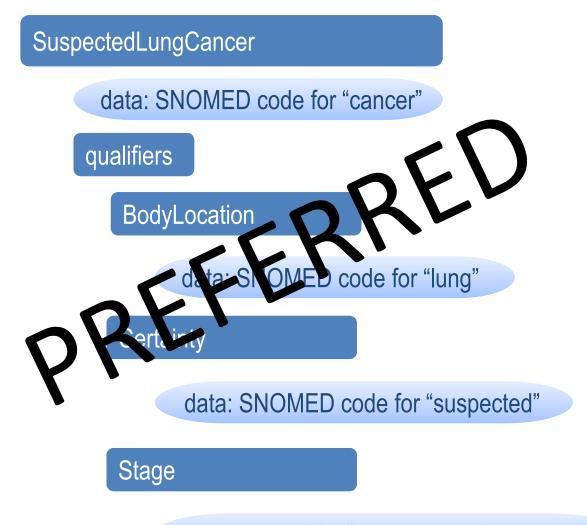
BodyLocation

data: SNOMED code for "lung"

Certainty

data: SNOMED code for "suspected"

Stage



SuspectedLungCancer

data: SNOMED code for "suspected cancer"

qualifiers

BodyLocation

data: SNOMED code for "lung"

Stage

SuspectedLungCancer

data: SNOMED code for "suspected lung cancer"

qualifiers

Stage

CIMI Repository

Objective: Create a single, shared repository of detailed clinical information models

- using approved formalisms
 - Archetype Definition Language (ADL)
 - Archetype Modeling Language (AML)
- with formal bindings to standard coded terminologies

Progress

- CIMI data types approved
- CIMI Reference Model approved
- A set of reference archetypes approved
- ~2000 CIMI lab models created
 - http://www.clinicalelement.com/cimi-browser/

Primary Near Term Goals

- Build a CIMI authoring tool
- As soon as possible, make more CIMI models available in a web accessible repository
- Get the models used in working systems
- Document our experience
- Improve our processes and models
- Repeat!



THE HEALTHCARE INNOVATION ECOSYSTEM

Wiki:

https://healthservices.atlassian.net/wiki/display/HSPC/Healthcare +Services+Platform+Consortium

Website: http://hspconsortium.org/#/

HSPC MISSION

Improve health by creating a vibrant, open ecosystem of interoperable applications

HSPC

- Provider-driven organization of healthcare organizations, IT vendors, systems integrators, and venture firms
- Incorporated as 501(c)(3) Aug 2014
 - -Chairman of the Board: Stan Huff, MD,Intermountain Healthcare/University of Utah
 - -CEO: Oscar Diaz, formerly VP, Harris Systems

HSPC Members To Date

Benefactor

- Intermountain Healthcare
- Louisiana State University
- Dept of Veterans Affairs

Associate

- Motive
- Regenstrief Institute

Individual

- American Medical Association
- Cognizant
- Jon Farmer
- Med Red
- Thomas Lang
- Regional Healthcare Improvement
- Wave Access

Sample of Participants

- HL7 FHIR Grahame Grieve
- SMART Josh Mandel
- Cerner David McCallie, Marc Overhage
- Epic Janet Campbell
- VA Jonathan Nebeker, Paul Nichol
- openEHR Thomas Beale
- Open Health Tools David Carlson
- Harris Vishal Agrawal
- Intermountain Healthcare
- Systems Made Simple Viet Nguyen
- LSU Frank Opelka, Wayne Wilbright, John Couk

- Center for Medical Interoperability Todd Cooper
- RelayHealth Arien Malec
- NLM Clem McDonald
- Infocare Healthcare Herb White
- Mayo Clinic Cris Ross
- Clinical Architecture Shaun Shakib
- Cognitive Medical Systems Doug Burke, Claude Nanjo, Emory Fry
- IBM Jeff Rogers, Dennis Leahy
- ASU Aziz Boxwalla, Robert Greenes
- Regenstrief Institute Douglas Martin
- U of Utah Ken Kawamoto

The **Essential** Functions

- Identify a set of services and their appropriate use cases
- Identify/create content (models/profiles) that reference standard terminology
- Identify protocols for security, authorization, context sharing, transport, etc.
- Publish the standards and development instructions openly, licensed free-for-use
- Provide conformance testing of software
- Engage vendors in supporting the standard services

Other Functions of the Consortium

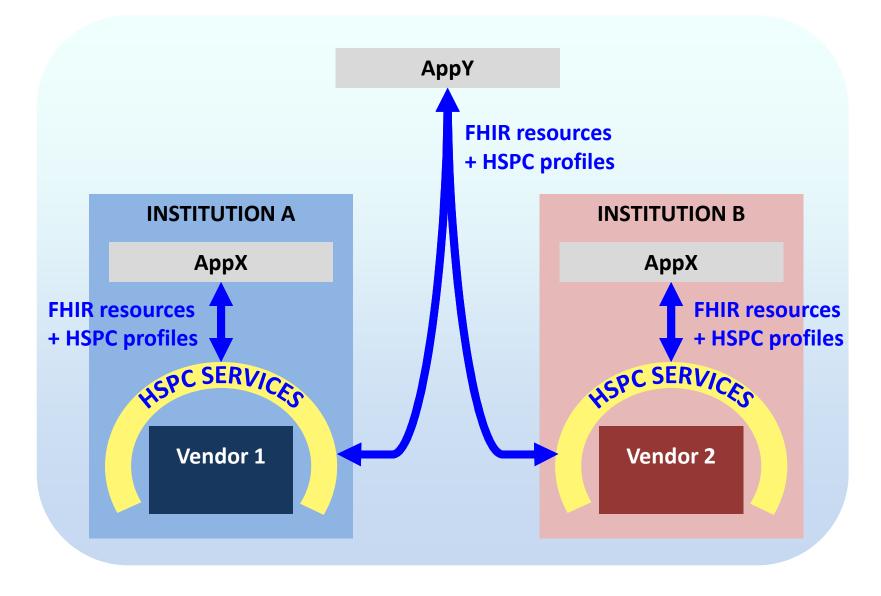
- Enable development "sandboxes"
- Set up a vendor- and provider-neutral "App Store"
- Create a business framework to support collaborative development
- Provide a way for people to invest (venture capital) in HSPC technologies

HSPC Technology Decisions

- Services FHIR
- Logical Data models (basis for FHIR profiles)
 - CIMI models + other existing content
- Terminology
 - LOINC, SNOMED CT, RxNorm, HL7 tables
- EHR Integration SMART on FHIR

How CIMI and HSPC other sources fit together **FHIR HSPC DSTU2** profiles Standard repository **Terminologies CIMI** repository **FHIR DSTU3** profiles CIMI **HTML** "Logical Model" CIMI **Translators Authoring Tool** models **RDF** HL7 v2 **FHIM** models HL7 v3 openEHR CDA Archetypes templates **CDISC SHARE**

The HSPC Mission

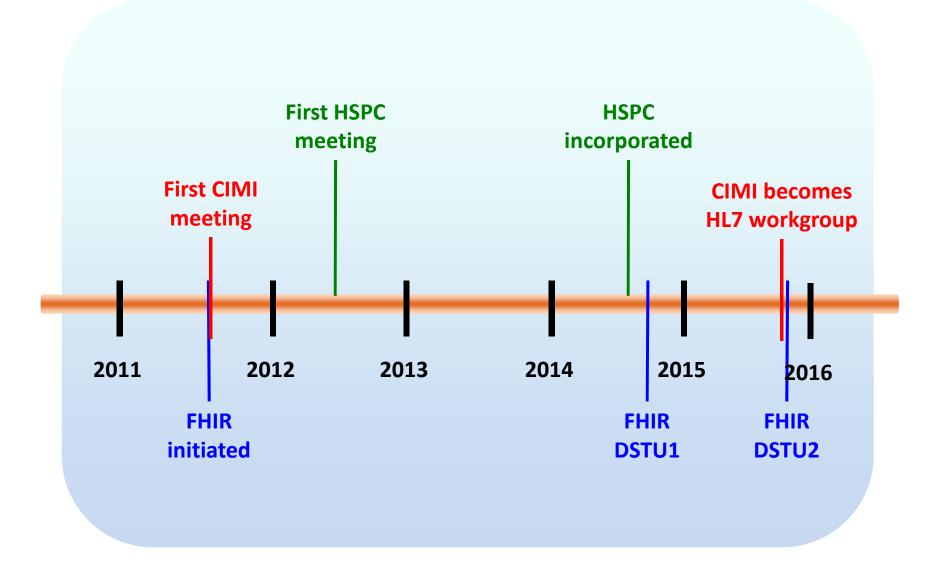


HSPC: More than just Data Virtualization

Specify standards for:

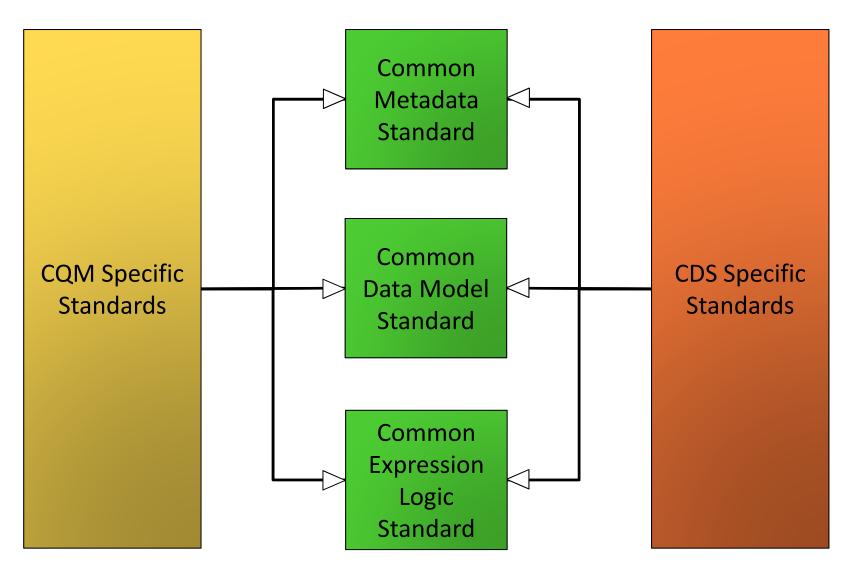
- Implementing a multi-layered services architecture (SOA)
- Supporting common Decision Support models (BPMN2/Drools)
- Supporting common workflow models (BPMN2)
- etc.

The HSPC, CIMI, and FHIR Timeline

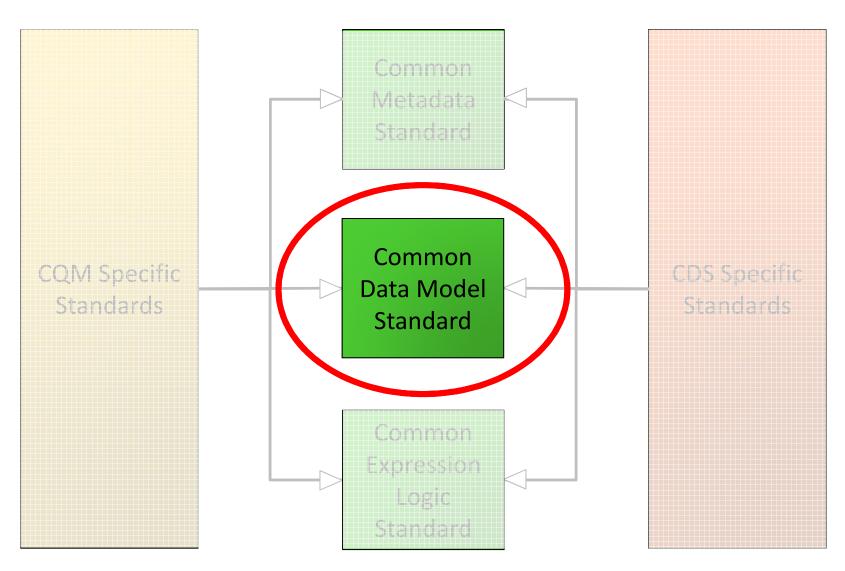


an evolving relationship . . .

CQF Value Statement (reproduced from wiki.siframework.org)



CQF Value Statement (reproduced from wiki.siframework.org)

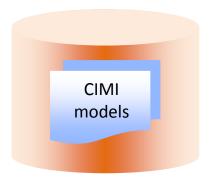


Why it Makes Sense

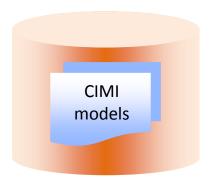
- Motivation for CIMI/HSPC is decision support and computable data
 - emphasis on coded data
 - tendency toward postcoordination
- CIMI already talking to modeling community
- Synergy -- CQF can leverage CIMI/HSPC experience and content and vice versa

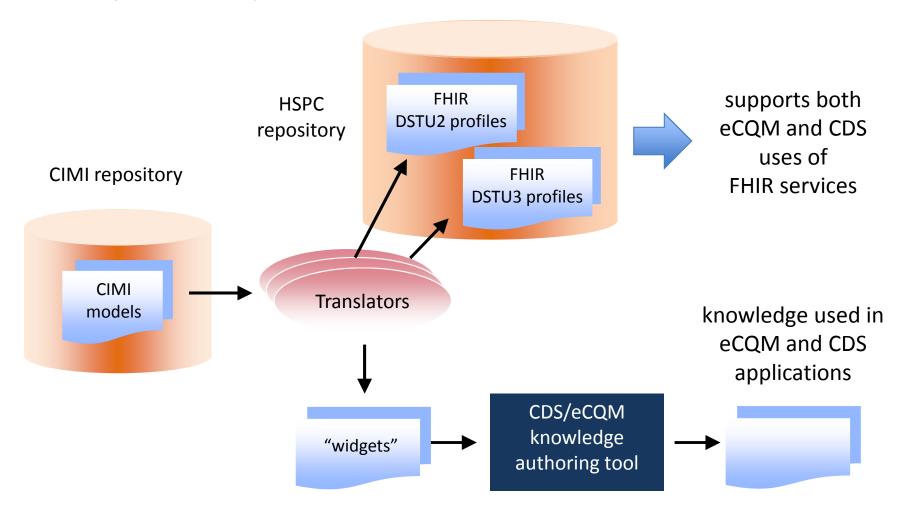
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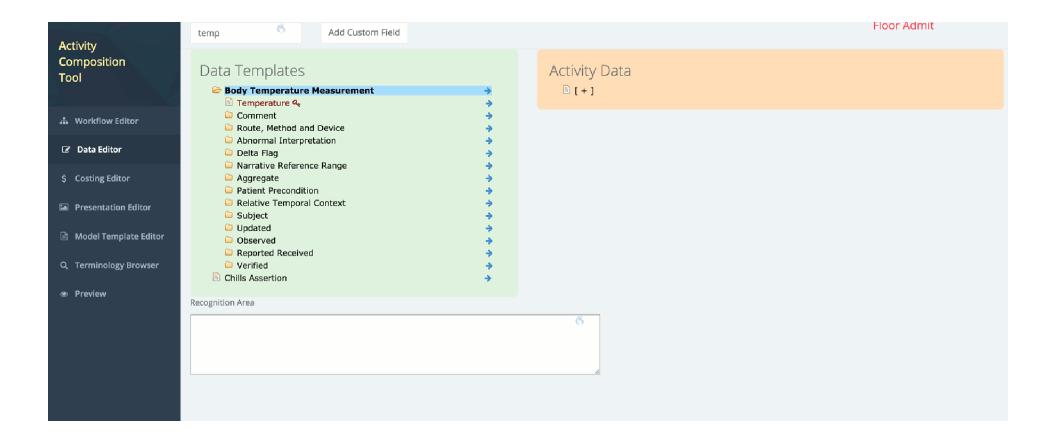
CIMI repository

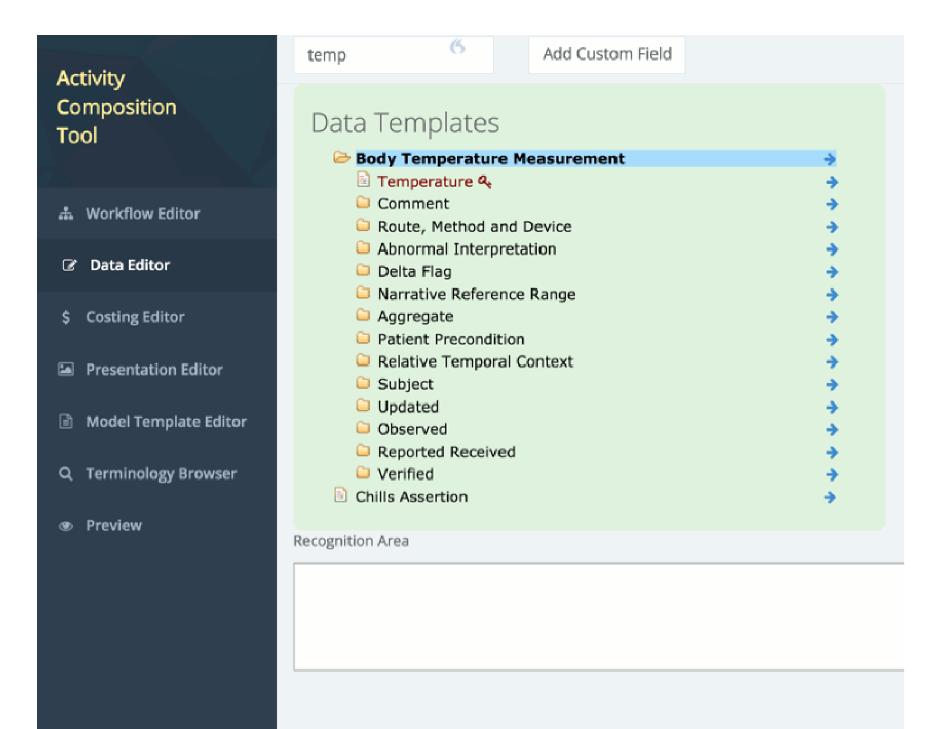


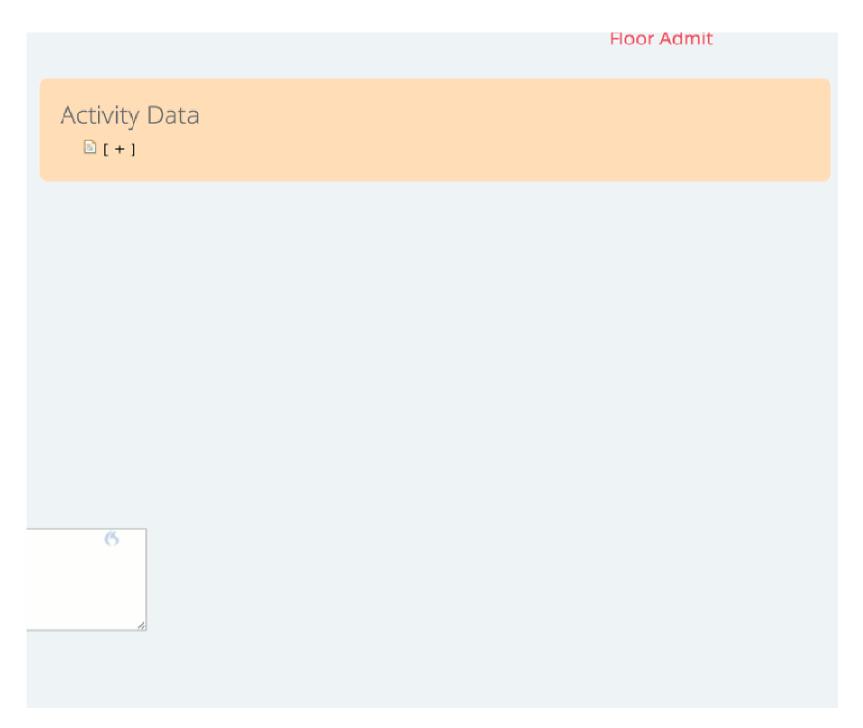
CIMI repository



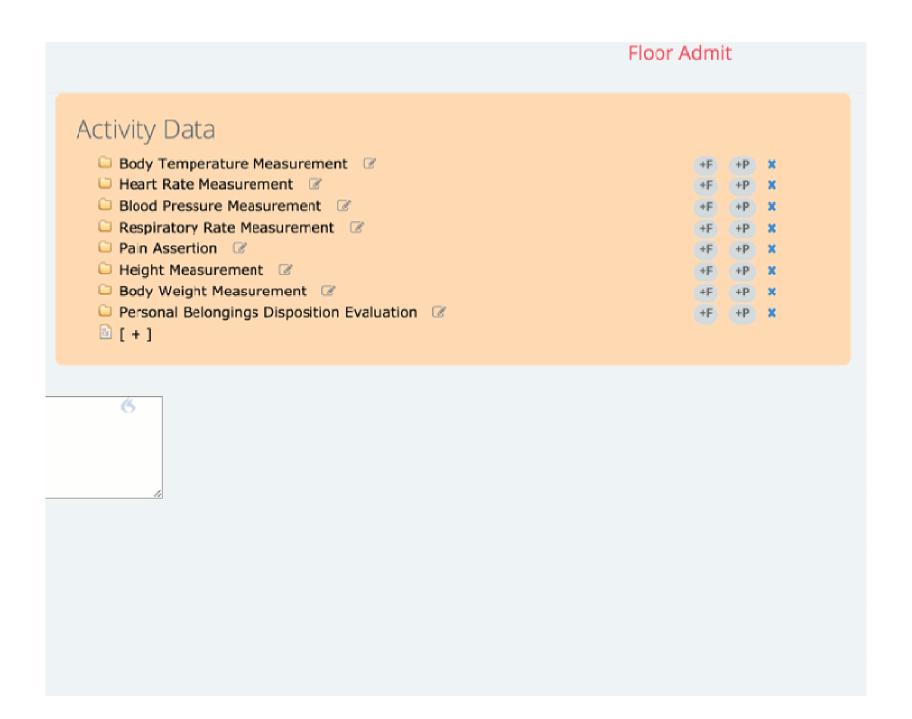








Floor Admit Activity Data Body Temperature Measurement [+]



Conclusion

- CIMI/HSPC can serve as the data model standard needed by the CQF effort.
- CIMI models can serve as the logical basis for HSPC FHIR profiles and CDS/eCQM knowledge artifacts adopted by CQF.
- CIMI/HSPC + CQF => synergy

DISCUSSION AND Q&A

Discussion Questions

- What do you think about the proliferation of standards and efforts to coordinate among standards development initiatives?
 Do you think the alignment efforts described are headed in the right direction?
- What recommendations do you have for standards development and implementation in this area?
- What gaps still exist in the available standards related to CDS and eCQM? How should we address those gaps?
- What clinical domain areas can most benefit from standardsbased, interoperable CDS and eCQM (e.g., immunizations, chronic disease management, chemotherapy)?
- How should CDS and eCQM be leveraged in the short-term and long-term to improve clinical quality and health outcomes?

Thank You!

| Panelist | Title | Initiative Role |
|--|--|--|
| Julia Skapik, MD, MPH Julia.skapik@hhs.gov | Medical Officer Office of the National Coordinator for Health IT | Executive Sponsor |
| Kensaku Kawamoto, MD, PhD, MHS kensaku.kawamoto@ utah.edu | Associate CMIO University of Utah | Co-Initiative Coordinator |
| Marc J. Hadley, PhD mhadley@mitre.org | Senior Principal Software Systems Engineer MITRE Corporation | Co-Initiative Coordinator |
| Tom Oniki, PhD Tom.Oniki@imail.org | Senior Medical Informaticist Intermountain Healthcare | Collaboration Coordinator, HSPC and CIMI |