

Monthly Summary Briefing HL7 EHR Work Group (EHR-WG)



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November 23, 2013Frequently-Updated Working-Drafthttp://wiki.hl7.org/index.php?title=EHR_Interoperability_WG



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The complete-and-current HL7 <u>EHR-System Function-and-Information Model Release-3</u> Development-Summary Presentation, dated November-2013 is available at <u>http://wiki.hl7.org/index.php?title=EHR_Interoperability_WG</u> 2



EHR Work Group Goal & Objectives

- <u>Electronic Health Record (EHR) Work Group's</u> goal is to support the HL7 mission of developing standards for EHR data, information, functionality, and interoperability.
 - Functional and Information Requirements for Electronic Health Records (EHR) and systems (EHRS),
 - Functional and Information Requirements for Personal Health Records (PHR) and systems (PHRS),
- EHR Interoperability WG's objectives are
 - 1. to create a clear, complete, concise, correct and consistent EHR-S FIM r3.0 in the Sparx Systems Enterprise Architect (EA) tool; where, it addresses the issues identified by the VA negative r2.0 ballot.
 - 2. to produce a Meaningful Use profile for r2.0.
- <u>Resource Management Evidentiary Support (RM-ES) project's</u> **objective** is to provide expertise on records management, compliance, and data/record integrity and governance to support the use of medical records for clinical care and decision-making, business, legal and disclosure purposes.
- <u>EHR Usability WG's</u> objective is developing a usability profile for the EHR-SFM
- <u>PHR-SWG's</u> objective is to maintain a Patient Healthcare System Functional Model (PHR-SFM).

EHR WG



Schedule:http://www.hl7.org/concalls/default.aspxList Server:http://www.hl7.org/myhl7/managelistservs.cfm

Health Level Seven – Electronic Health Record Work Group Weekly Teleconference Schedule

Revised: 20 November 2013

Day	Time US ET	Activity	Lead(s)	Dial-In	Screen Sharing	List Server (for agendas, announcements)		
Mon	1200	Records Management/ Evidentiary Support	Warner, Gelzer	1-877-668-4493 Code 927 002 088#	<u>Link</u>	EHR Legal		
Tues	1300	EHRS FM Release 3 Planning	Hufnagel, Dickinson	1-770-657-9270, Passcode 510269#	<u>Link</u>	EHR Interop		
	1400	Meaningful Use Functional Profile	Datta, Dickinson	1-770-657-9270, Passcode 510269#	<u>Link</u>	EHR Interop		
	1500	FULL EHR WG	Co-Chairs	1-770-657-9270, Passcode 510269#	<u>Link</u>	EHR WG		
Wed	1200	Personal Health Record WG	Ritter, Dickinson, Doo	1-770-657-9270, Passcode 510269#	TBA	EHR PHR		
	1300	EHR System Usability WG	Mon, Ritter, Rocca, Gartner	1-770-657-9270, Passcode 510269#	<u>Link</u>	EHR Usability		
Thur	Open							
Fri	0930	EHR WG Co-Chairs	Co-Chairs	1-770-657-9270, Passcode 510269#	TBA	N/A		

EHR-S FIM Acronyms

also known as aka . CC EHR-S FIM Conformance Criteria ٠ CDA **Clinical Document Architecture** ٠ DD **Data Dictionary** ٠ CIM **Conceptual Information Model** ٠ СР Care Provision ٠ CPS Care Provisioning Support ٠ EA Enterprise Architect ٠ EHR-S EHR System ٠ EHR-S FIM EHR-S Function and Information Model ٠ FHA US Federal Health Architecture ٠ FHIM US Federal Health Information Model ٠ Fast Healthcare Interoperability Resources FHIR ٠ EHR-S Function and Information Model FIM ٠ FIM(MU) EHR-S FIM profile for MU ٠ **Function Model** FM ٠ FY **Fiscal Year** ٠ IM Information Model ٠ MDHT Model Driven Health Tools ٠ MU US Meaningful Use objectives-and-criteria ٠ ONC US Office of the National-Coordinator ٠ **Open Health Tools** OHT ٠ **POA&M** Plan of Actions and Milestones ٠ R 2/3 Release 2 or 3 ٠ RI Resource Infrastructure ٠ HL7 Reference Information Model RIM ٠ S&I ONC Standards & Interoperability Framework ٠ WBS Work Breakdown Structure ٠ WG Work Group .

Executive Summary EHR-S FIM r3:2016 Preparation



This executive-summary specifically addresses potential work-group impacts and/or trends, which are important for VA, IPO and DOD awareness.

EHR System Function-and-Information Model (EHR-S FIM)

• Structured, based-on a fully-specified Reference Model (RM) for

- Clear, complete, concise, correct, consistent and intuitive ease-of-use;
- Sparx Enterprise Architect (EA) UML-model tool-based; where, release 3 (r3)
 - manages user-activities, system-functions. business-rules, interoperable-data separately; and,
 - Consistent-global Conformance Criteria (CCs) replace ad-hoc-local r2 CCs
 - Single Infrastructure-section contains previously-separate r2 Record-and-Trust Infrastructure-sections
- EA Tool-generated Interoperability-Specifications based-on Use-Cases
 - <u>Use-Cases</u> come-from HITSP & S&I Framework Use-Case Simplification work linked-to
 - <u>Requirements</u>, which come-from EHR-S r2.0 Functions' and their restructured CCs linked-to
 - International Interoperability-Specifications based-on HL7 FHIR (Fast Healthcare Interoperability Resources)
 - <u>US-Realm</u> Interoperability-Specifications based-on FHAFHIM (Federal Health Information Model)

NOTE: EHR-S FIM is <u>NOT</u> intended to imply a specific architecture or workflow!

Executive Summary Conclusions and Recommendations EHR-S FIM r3:2016 Preparation



- 1. EHR-S FIM vision is to become the <u>"Easy Button" for EHR Interoperability Specifications</u>
 - a. Easily-customizable to user-specific profiles.
 - b. Including a US-Realm Meaningful Use (MU) & FHIM profile
 - c. EHR-S FIM r3:2016 within Sparx EA represents a powerful HL7 product; where,
 - i. EA integrates FHIR, FHIM and S&I Framework's Use-Case Simplification, and
 - ii. The EA tool-based EHR-S FIM is Governed and Configuration Managed consistently.
 - iii. The EA tool can generate both a navigable-web-site and printable-report
 - iv. Support user-specific profiles (e.g., WG project DAMs, DIMs, DCMs).
- 2. HL7.org/EHRSFIM web-site should be setup-and-managed by the EHR Interoperability WG
 - a. Supporting peer review, trial-use and stakeholder-contribution during FY14-Alpha & FY15-Beta development.
- 3. EHR-S FIM development, tooling and balloting resources = (estimated) 5-FTE Man-years a. A marketing campaign is needed to justify EHR-S FIM r3:2016 resources

Plan of Actions and Milestones FY2014Q1 POA&M EHR-S FIM Release-3:2016 Preparation						
October 2013 (Identify processes, tools and issues/risks)	Completed					
Prototype CP.6.2 Immunization Management	22-Oct-13					
 Prototype RI.1.1.1 Originate and Retain Record Entry 	29-Oct-13					
November 2013 (Prototype complete process-and-products)						
Prototype FHIR integration (Allergies, Intolerance & Adverse Reaction)	5-Nov-13					
Prototype FHIM integration (Allergies, Intolerance & Adverse Reaction)	8-Nov-13					
 Define EHR-S Reference-Model and Conceptual-Architecture 	15-Nov-13					
Prototype Use-Case generation of Immunization Interoperability-Specification	in-progress					
 Harmonize with ISO/EN 13940 Continuity-of-Care System-of-Concepts 	pending					
Harmonize with Electronic Health Record Communication (ISO/EN 13606)						
 Prototype EHR-S FIM Ballot Production process-and-products for prototype 						
December 2013 (Develop production WBS and POA&M)						
 Create Release 3 Work-Break-Down Structure (WBS) & POA&M 						
Setup EA tool with finalized Release 2, after ISO ballot reconciliation						
January 2014 – 2016 (Approve & Execute Plan)						
• Jan 2013: Present Prototype, WBS & POA&M at HL7 WG meeting; then, execute	e POA&M.					

• Establish public <u>www.EHR-S-FIM.org</u> website to get broad peer-review

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Concept-of-Operations Scenario EHR-S FIM Release-3:2016 Preparation

A <u>Clinician</u> and <u>Patient</u> and/or their designated <u>Agents</u> have <u>Encounters</u>; where, they may use <u>EHR-S</u> (EHR System) <u>GUI</u> (Graphical-User-Interface) to *manage* Data, in accordance with scope-ofpractice, organizational-policy, jurisdictional-law, and patient-preferences:

- The <u>Clinician</u> reviews the <u>Patient</u>'s <u>EMR</u> (Electronic Medical Record) and related <u>Information</u>
- The <u>Clinician</u> observes, treats, writes-orders and documents the <u>Patient-Encounter</u>
- The <u>Patient</u> provides requested-<u>Information</u> and is provided educational-<u>Information</u>
- The EHR-S Manager manages
 - <u>Record Entries</u> for <u>Orders</u>, <u>Treatments</u> and <u>Observations</u>
 - <u>EMR Care-Plans</u>, <u>Care-Records</u>, and <u>Problems-and-Concerns</u>, which are organized from the <u>Record-Entries</u>.
 - Encounters, which are signed by the Clinician (s) and possibly the Patient.

Concept of Operations Model EHR-S FIM Release-3:2016 Preparation

uc EHR-S Concept-of-Operation



INTERNATION

Manager Operations aka Verb Hierarchy EHR-S FIM Release-3:2016 Preparation

uc EHR-S Manager Anatomy (Verb Hierarchy)



INTERNATIONA



Reference Model Definition EHR-S FIM Release-3:2016 Preparation

The EHR-S reference model (RM) is an abstract-framework for structuring significant-relationships among the entities of EHR-S environments basedon consistent EHR-S function-and-information conceptual models; where, EHR-S RM conformance criteria contain a constrained-lexicon of nouns (entities), verbs (operations/tasks), qualifiers (conditions), constraints (policies/rules), which may be used-as requirements-specifications by analysts, developers, implementers, and testers. The EHR-S or PHR-S RMinstance-models provide a common syntax-and-semantics that can be used unambiguously across-and-between different implementations; where, the may be linked-to specific-implementation standards-RM instances technologies-paradigms-or-patterns. [based-on OASIS RM definition]

[•] According to the Organization for the Advancement of Structured Information Standards (**OASIS**) a reference model is "an abstract framework for understanding significant relationships among the entities of some environment, and for the development of consistent standards or specifications supporting that environment. A reference model is based on a small number of unifying concepts and may be used as a basis for education and explaining standards to a non-specialist. A reference model is not directly tied to any standards, technologies or other concrete implementation details, but it does seek to provide a common semantics that can be used unambiguously across and between different implementations."

EHR-S RM Conformance-Criteria Definition EHR-S FIM Release-3:2016 Preparation



• Human

- Clinician, Patient, Designated Agent

• Human Action

- Linked-to Use-Case Actions
- CLINICIAN: Observe, Treat, Write-Orders
- such as Immunization Administration
- System
 - EHR or PHR
- Applicability (SHALL, SHOULD or MAY)
 - according to
 - Scope of practice,
 - Organizational policy,
 - Jurisdictional law,
 - Patient preference or consent."

• System Function Constraints

- Invariant-conditions (e.g., context)
- Pre-conditions (e.g., triggers)
- Post- conditions (e.g., goal, outcomes)

• System Function Type

- System provides the ability (for a human) to
- Or the system directly does

System Function

EHR Verb Hierarchy of what the system does, such as manage, maintain, ...

• Data Requirements

- Linked-to International FHIR specifications
- Linked-to US Realm FHIM specifications

Associations & Dependencies

- Supporting capabilities and functions

EHR-S RM Example Conformance-Criteria EHR-S FIM Release-3:2016 Preparation

CP.6.2#01 <u>During an Encounter</u>, the EHR system SHALL provide the ability to *capture* <u>Immunization Administration</u> details as discrete data, such as Immunization FHIR (Fast Healthcare Interoperability Resource); where, the Immunization data may be associated with the following resources:

- AdverseReaction and other Observations;
- Patient , Practitioner, Organization, Location

And, within the US Realm, the Immunization and associated resources are express-able by the applicable FHIM (Federal Health Information Model) Domains of:

- Immunization, Adverse Reaction, Allergy and Intolerance
- Associated with appropriate FHIM classes (e.g., Person, ...)

Use-Case Traceability Analysis CP.6.2 Immunization Management Conformance Criteria



IC EHR-S FIM CP.6.2 Immunization Management Traceability



Example Linkage-to FHIR & FHIM for Allergy, Intolerance & Adverse-Reaction



class FHIM Allergy, Intolerance and Adverse Reaction



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Use-Case Description CP.6.2 Immunization Management EHR-S FIM Release-3:2016 Preparation



- A <u>Clinician</u> manages the EHR-S to
 - reviews the patient's <u>EMR</u> for <u>Allergies and Intolerances</u>, <u>Immunization-Schedule</u>, and <u>Patient Directives</u>
 - Document treatment (*immunizes*) of the <u>Patient</u> with a <u>Vaccine</u> and <u>Adverse-Reaction</u> observations.
 - documents the encounter information.
- The EHR-S Immunization related manager(s) can
 - Capture, Auto-populate, Maintain, Render, Transmit, Exchange,
 - Harmonize, Update, or Determine
- The following data-modules:
 - Immunization-Administrations, Allergies, Intolerances, Adverse-Events
 - Events, Schedules, Plans and Educational Materials

Use-Case Model CP.6.2 Immunization Management Conformance Criter

c EHR-S FIM CP.6.2 Immunization Management (Clinician View)



SHALL, SHOULD or MAY applicability is "according to scope-of-practice, organizational-policy, jurisdictional-law, patient preference-or-consent."





EHR-S-FIM Traceability Model CP.6.2 Immunization Management



class EHR-S FIM CP.6.2 Immunization Management (Conceptual Traceability Model)



EHR-S FIM Logical Traceability-Model CP.6.2 Immunization Management

class EHR-S FIM CP.6.2 Immunization Management (Logical Model)



EHR-S FIM Logical Traceability-Model CP.6.2 Immunization Management

class EHR-S FIM CP.6.2 Immunization Management (Logical Model-2)



EHR-S-FIM Conformance Criteria (CCs) CP.6.2 Immunization Management

- 1. The system SHALL provide the ability to capture, maintain and render immunization administration details as discrete data, including:(1) the immunization name/type, strength and dose;(2) date and time of administration;(3) manufacturer, lot number, expiration date,(4) route and site of administration;(5) administering provider;(6) observations, reactions and complications;(7) reason immunization not given and/or immunization related activity not performed; according to scope of practice, organizational policy and/or jurisdictional law."
- 2. The system MAY *auto-populate the immunization administration record* as a by-product of verification of administering provider, patient, medication, dose, route and time according to scope of practice, organizational policy and/or jurisdictional law.
- 3. The system **SHALL** provide the ability to *determine and render required immunizations*, and when they are due, based on widely accepted immunization schedules, when rendering encounter information.
- 4. The system SHOULD provide the ability to capture, in a discrete field, an allergy/adverse reaction to a specific immunization.
- 5. The system **SHALL** conform to function CP.3.2 (Manage Patient Clinical Measurements) to capture other clinical data pertinent to the immunization administration (e.g., vital signs).
- 6. The system SHOULD provide the ability to link standard codes (e.g. NDC, LOINC, SNOMED or CPT) with discrete data elements associated with an immunization.
- 7. The system **SHALL** provide the ability to *maintain the immunization schedule*.
- 8. The system **SHALL** provide the ability to render a patient's immunization history upon request for appropriate authorities such as schools or day-care centers.
- 9. The system **SHALL** conform to function CP.1.2 (Manage Allergy, Intolerance and Adverse Reaction List).
- 10. The system SHOULD transmit required immunization administration information to a public health immunization registry according to scope of practice, organizational policy and/or jurisdictional law.
- 11. The system SHOULD exchange immunization histories with public health immunization registries according to scope of practice, organizational policy and/or jurisdictional law.

EHR-S-FIM Conformance Criteria (CCs) CP.6.2 Immunization Management



ISSUE: Consistency of Conformance Criteria (CC) across related functions, such as Medication-and-Immunization and Orders-and-Results Management.

- 12. The system SHOULD harmonize Immunization histories with a public health immunization registry according to scope of practice, organizational policy and/or jurisdictional law.
- 13. The system SHOULD capture and render immunization histories from a public health immunization registry.
- 14. The system SHALL conform to function CP.1.6 (Manage Immunization List).
- 15. The system SHOULD provide the ability to update immunization histories at the time of capturing an immunization administration.
- 16. The system **SHALL** provide the ability to render the immunization order as written (i.e., exact clinician order language) when rendering administration information.
- 17. "The system SHALL provide the ability to determine due and overdue ordered immunizations and render a notification. "
- 18. The system **SHALL** provide the ability to render a patient educational information regarding the administration (e.g., Vaccine Information Statement (**VIS**)).
- 19. The system **SHALL** provide the ability to capture that patient educational information (e.g., VIS) was provided at the time of immunization administration.
- 20. The system **SHALL** provide the ability to capture documentation that patient educational information (e.g., VIS) was provided at the time of immunization administration.
- 21. The system **SHALL** provide the ability to capture the receiving entity (e.g., patient, representative, organization) when patient education information is provided at the time of immunization administration.
- 22. The system SHOULD provide the ability to capture and maintain immunization refusal reasons as discrete data.
- 23. The system SHOULD provide the ability to capture patient preferences regarding receipt of immunization (e.g. refusal of certain vaccine types) at time of immunization administration.



Interim Conclusions EHR-S FIM r3.0:2016

- We have looked at Medication-and-Immunization Management, Orders-and-Results Management and Record Entry Management.
- The <u>EHR-S RM (reference model)</u> is used to structure EHR-S functions-and-data; where, the function's conformance-criteria lexicon defines the grammar of nouns (entities), verbs (record-entry actions) and constraints (conditions).
- The EHR-S <u>Conceptual Information Model (CIM)</u> and <u>Conceptual Operations Model (COM)</u> for CP.6.2 Immunization Management should generally-be-applicable for all of the Care Provisioning (CP) section of the EHR-S FM; where,
 - minor CIM modifications will likely occur as we analyze the rest of the CP section; but,
 - major COM components still must be substantially developed based-on the Record-Infrastructure and Trust-Infrastructure sections.

Interim Conclusion EHR-S FIM CP.6.2 Immunization Management



- Based on the Medication Management, Orders Management and Immunization Management functions, we see
 - A high-level EHR-S Information Model emerging as a set of
 - Patients, Providers, External Partners, Encounters, EMRs, Care Plans, Lists, Managers, Documents and Notes;
 - A high-level EHR-S Manager Model is emerging to
 - Capture, Auto-populate, Maintain, Render, Transmit, Exchange, Harmonize, Update, Determine

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EHR-S FIM Conceptual Information Model (CIM) RI.1.1 Originate and Retain Record Entry

class RI.1.1.1 Originate and Retain Record Entry (Conceptual Traceability View)



INTERNATION

EHR-S FIM Traceability View RI.1.1.1 Originate-and-Retain Record Entry







Conformance Criteria (CC) RI.1.1.1 Originate-and-Retain Record-Entry

- 1. RI.1.1.1#01 The system **SHALL** provide the ability to capture (originate) a Record Entry instance corresponding to an Action instance and context.
- 2. RI.1.1.1#02 The system SHALL capture a unique instance identifier for each Record Entry.
- 3. RI.1.1.1#03 The system **SHALL** conform to <u>function TI.2.1.1.1</u> (Originate/Retain Record Entry Audit Trigger), including specified metadata.
- 4. RI.1.1.1#04 The system **SHALL** capture the signature event (e.g., digital signature) of the origination entry Author, binding signature to Record Entry content.
- 5. RI.1.1.1#05 The system **SHALL** provide the ability to capture both structured and unstructured content in Record Entries.
- 6. RI.1.1.1#06 The system SHALL provide the ability to capture Record Entries from information recorded during system downtime.
- 7. RI.1.1.1#07 The system SHOULD provide the ability to integrate Record Entries from Information recorded during system downtime.
- 8. RI.1.1.1#08 The system **SHALL** provide the ability to capture date/time an Action was taken or data was collected if different than date/time of the Record Entry.
- 9. RI.1.1.1#09 The system SHOULD capture metadata that identifies the source of non-originated Record Entry (e.g., templated, copied, duplicated, or boilerplate information).
- 10. RI.1.1.1#10 The system MAY provide the ability to tag unstructured Record Entry content to organize it according to need, for example, in a time-related fashion or by application-specific groups (such as photographs, handwritten notes, or auditory sounds)
- 11. RI.1.1.1#11 The system MAY capture and maintain a Record Entry encoded as a standards-based data object (e.g., HL7 Continuity of Care or other HL7 CDA R2 Document).
- 12. RI.1.1.1#12 The system MAY capture and maintain a standards-based data object to mirror (be duplicate and synchronous with) internal Record Entry representation.

EHR-S FIM Logical View RI.1.1.1 Originate-and-Retain Record Entry

class RI.1.1.1 Originate and Retain Record Entry (Logical View)



EHR-S FIM



RI.1.1.1 Originate and Retain Record Entry

Resultant Description (Notional Scenario)

- The EHR-S <u>Record-Entry</u> manager can
 - Capture, Create, Copy, Record, Transcribe, Identify,
 - Link, Tag, Encode, Mirror, and Integrate
- <u>Record-Entries</u> as
 - structured or unstructured-data link-to associated
 - Event-Metadata and Signatures.



we have only looked at the RI.1.1.1 function; yet,

- we see that the emergence of common <u>Record-Entries</u>, <u>Events</u>, <u>Record Entries</u> and a <u>Record Entry Manager</u>
- which can Capture, Create, Copy, Record, Transcribe, Identify, Link, Tag, Encode, Mirror, Integrate
 - structured-data or unstructured-data and link-to
 - associated Event-Metadata and Signature.

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EHR-S FIM Using FHIR

ISSUE: EHR-S FM r2.0 Implied Information Model is Ad-Hoc; where, FHIR & FHIM Information Model & Data Dictionary are Configuration Managed.

FHIR Administrative

- Attribution: Patient, RelatedPerson, Practitioner, Organization
- Resources: Device, Location, Substance, Group
- Workflow Management: Encounter, Alert, Supply, Order, OrderResponse
- Financial: Coverage

FHIR Clinical

- General: AdverseReaction, AllergyIntolerance, CarePlan, FamilyHistory, Condition, Procedure, Questionnaire
- Medications: Medication, MedicationPrescription, MedicationAdministration, MedicationDispense, MedicationStatement, Immunization, ImmunizationProfile
- **Diagnostic**: Observation, DiagnosticReport, DiagnosticOrder, ImagingStudy, Specimen
- Device Interaction: DeviceCapabilities, DeviceLog, DeviceObservation

FHIR Infrastructure

- **Support**: List, Media, Other, DocumentReference, (Binary)
- Audit: Provenance, SecurityEvent
- **Exchange**: Document, Message, OperationOutcome, Query
- Conformance: Conformance, ValueSet, Profile

EHR-S FIM Prototype Allergy, Intolerance & Adverse-Reaction FIM-FHIR-FHIM Requirements-Specifications ISSUE: Should we map at Data Module Level or Conformance Criteria level? [Gary]



Prototype Allergy, Intolerance & Adverse-Reaction FHIR Design-Specification



class FHIR Specification for Allergy, Intolerance and Adverse Reaction



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EHR-S FIM Using Federal Health Information Model (FHIM) http://www.fhims.org/content/420A62FD03B6_root.html



FHA Federal Health Information Model (FHIM)



Prototype Allergy, Intolerance & Adverse-Reaction FHIM High-Level US-Realm Specification

class FHIM Allergy, Intolerance and Adverse Reaction



Prototype FHIM-Detailed Allergy & Intolerance Specification

class FHIM Allergies Domain





Prototype FHIM Detailed Adverse-Reaction Specification

class FHIM Adverse-Event Reporting Domain



Prototype Allergy, Intolerance & Adverse-Reaction FHIR & FHIM Design-Specifications INTERIM CONCLUSION

- EHR-S FIM, FHIR and FHIM complement each other; where,
- EHR-S FIM defines <u>Requirements</u>; where,
 - EHR-S FIM needs data-specifications and Dictionary and
 - FHIR & FHIM provide data-specifications and Dictionary
- FHIR defines the International Data-Specifications ("The 80% set")
- FHIM can define the US-FHA FHIR-Profile
- Joint Configuration Management is essential for FIM/FHIR/FHIM consistent
- A FIM-FHIR-FHIM populated UML-Tool (e.g., EA or RSA) can manage
- **Requirements** from EHR-S FIM
- International Data-Specifications from FHIR
- US-Realm Data-Specifications-Profile from FHIM

Contents FY2014Q1-Prototype Report EHR-S FIM Release-3:2016 Preparation

- 1. Introduction, Executive-Summary, Plan-of-Actions & Milestones
- 2. Concept-of-Operation and Reference-Model
- 3. CP.6.2 Immunization-Management Modeling-Prototype
- 4. RI.1.1.1 Originate and Retain Record Entry Modeling-Prototype
- 5. EHR-S FIM use of FHIR for Allergy, Intolerance and Adverse-Reaction
- 6. EHR-S FIM use of FHIM for Allergy, Intolerance and Adverse-Reaction
- 7. Traceability

The complete-and-current HL7 <u>EHR-System Function-and-Information Model Release-3</u> Development-Summary Presentation, dated November-2013 is available at <u>http://wiki.hl7.org/index.php?title=EHR_Interoperability_WG</u> 46



EHR-S FIM Issue Traceability

ISSUE: EHR-S FM r2.0 traceability to UML Model Elements to EHR-S FIM r3.0, FHIR & FHIM

- Workbook 1: Class attributes & operations mapped-to EHR-S FM r2.0 Functions and LOCALCCs
- Workbook 2 Class attributes & operations mapped to EHR-S FIM r3.0 Functions and GLOBALCCs
- Workbook 3 EHR-S FM r2.0 Functions and CCs mapped-to EHR-S FIM r3.0 Functions and CCs
- Workbook 4 EHR-S FM r2.0 Functions and LOCAL Conformance Criteria (CC) listed out for linking
- Workbook 5 EHR-S FIM R3.0 Functions and UNIVERSALCC listed out for linking
- Workbook 6 EHR-S FIM UML-Model mapped-to FHIR
- Workbook 7 EHR-S FIM UML-Model mapped-to FHIM (Federal Health Information Model)
- Workbook 8 FHIR mapped-to FHIM (Federal Health Information Model)
- Workbook 9 Master Data Dictionary (DD) (If we use FHIR or FHIM, they already have a DD)
- **ACTION:** Use Sparx EA to implement t raceability.

EHR-S FM Action-Verb Hierarchy Vs. EHR-S FIM Manager-Operations VS. Record Lifecycle Events



ISSUE: traceability of CC Verb-Hierarchy vs. Record Lifecycle Events.

Manage (Data)

Capture	Maintain			Render			Exchange	Determine		Manage- Data- Visibility
Auto- Populate Enter Import Receive	Store Archive Backup	Update Annotate	Remove Delete	Extract Present Train Record Entry Lifecycle Event Type Enumeration - originate and retain - originate and retain - amend - translate - attest - output/report - disclose - transmit - receive and retain - de-identify - pseudomynize - re-identify - extract - archive	Transr	smit	Export Import Receive Transmit	Analyze	Decide	De-Identify Hide Mask Re-Identify
	Decrypt Encrypt Recover Restore Save	Edit Harmonize Integrate Link Tag	Purge		ate ate t/report se hit e and retain entify omynize ntify t e e e (previously an	tain				
				 de stro de prev re-acti merge unmer link unlink place release 	y or identify m ciate/retract vate ge con legal hold e from legal hold	issing	← are but Ve	← Record-E are located but, how do Verbs in the	Entry Lifecycle Events here for convenience; they correspond to verbs hierarchy?	