

Electronic Record Services B.V.

HSE ISF ERS proposal

ERS

9 April 2014



Introduction: ISF



Irish National ISF

National ISF Core Objective

"A Standards Based Framework which will ensure key systems and components, can share data in timely and organised fashion."



Irish National ISF

National ISF - Key Strategic Objectives:

- Cost Avoidance
- Greater Efficiency
- Better Control
- 3rd party delivery of systems
- To facilitate the development of Electronic Health Records (EHR)
- To enable horizontal data integration and overcome data silos
- To facilitate a more controlled development environment
- Greater re-usability of key components



Irish National ISF

Approach

- Approach based on 12 Work streams
- Suppliers Framework for provision of specialist services
- LOT: 4 Tender for HSE National Information Architecture Model
- Award made to ERS B.V. in the Netherlands
- LOT 4: Now at final draft stage



Irish National ISF LOT 4

- Simplified overview of the model and key components
- Understanding the recommendations
- Making it real and deployable in simple terms
- Next stage 'quick win' options
- Simple deployment scenarios
- Costs and Benefits



HSE ISF IA-RM and SAM

ERS advocates a dual model approach as determined by the consultations and the Proof of Concept initiatives

(I) Information Architecture Reference Model

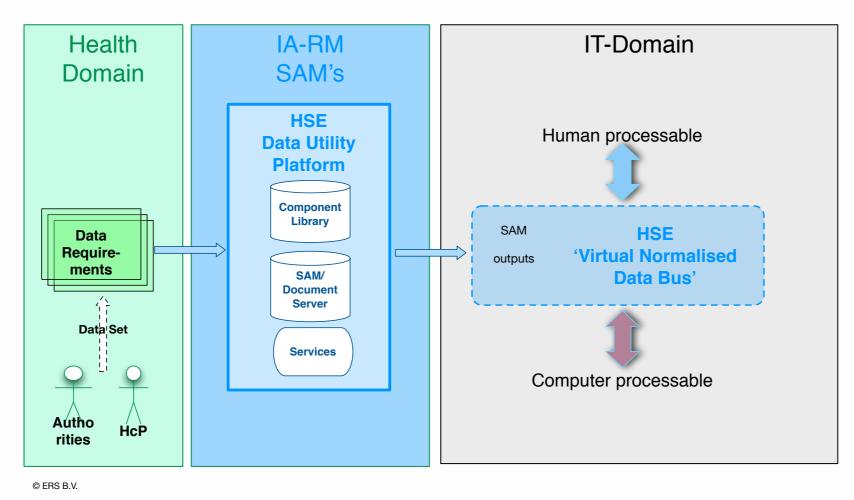
IA-RM as the basis for deployment in Ireland

(2) Subject Area Models

 SAM's as the expression of consensus data sets needed for Healthcare Providers to collaborate in clinical pathways





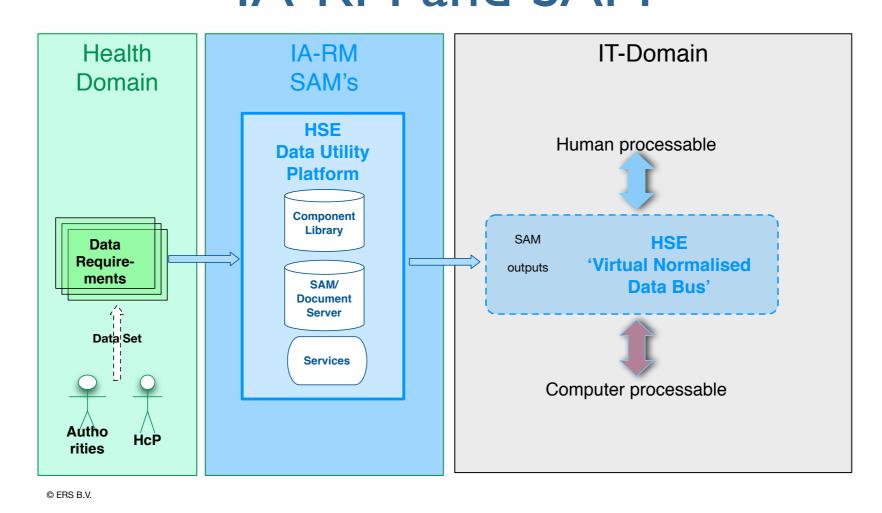


Data requirements are transposed to a data set (e.g. as defined by a clinical programme)

The Data Set is then transposed to a Subject Area Model







A series of tools and services (the HSE Data Utility Platform) is needed to generate SAM's

From SAM's different outputs (human readable, technical) are generated These are needed by all actors (Health- and IT-Domains).

When the SAM's are deployed via Procurement a 'Virtual Normalised Data Bus' will create a level green playing field for IT-vendors and other actors, based on open International standards



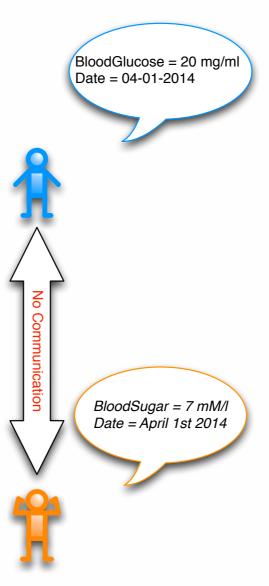
Information Architecture Reference Model

ERS

6 April 2014





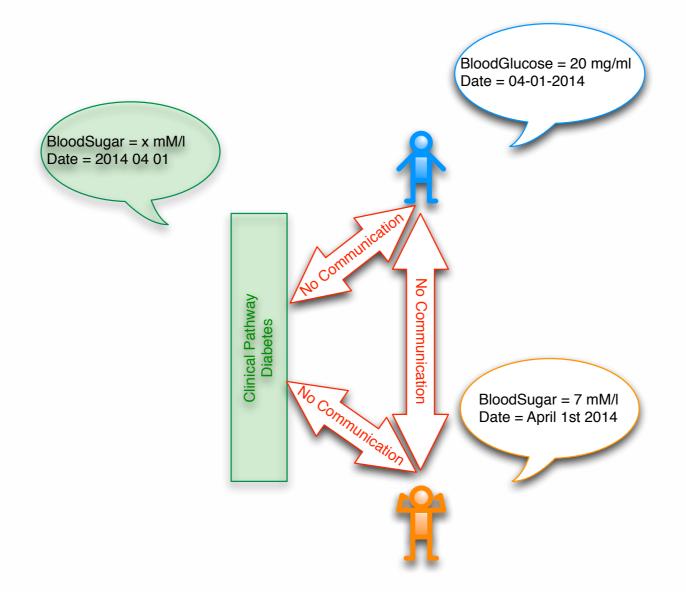


Why is an Information Architecture necessary?

- Both collaborating actors need to 'understand' each other
- Humans know how to bridge the gap; Computers do not
- Machine exchange of data needs very detailed and precise uniform agreements between IT-systems



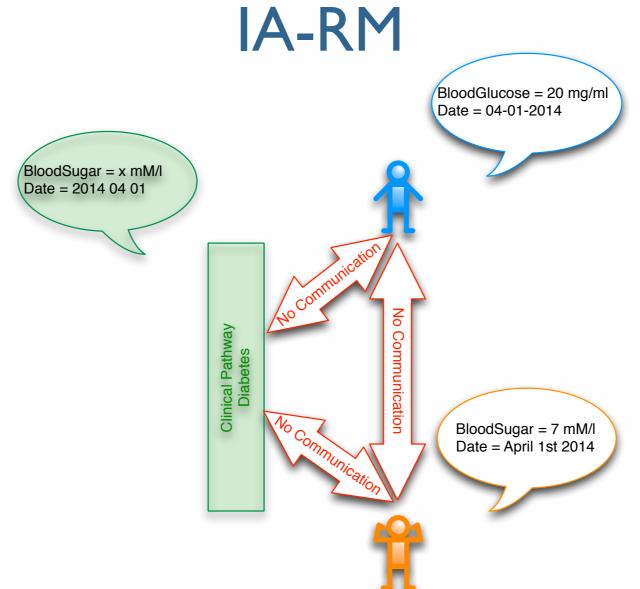




- The Information Architecture Reference Model (IA-RM) describes how collaborating actors and their IT-systems must exchange data
- Actors need to adapt to all the standards defined in the IA-RM

IA-RM



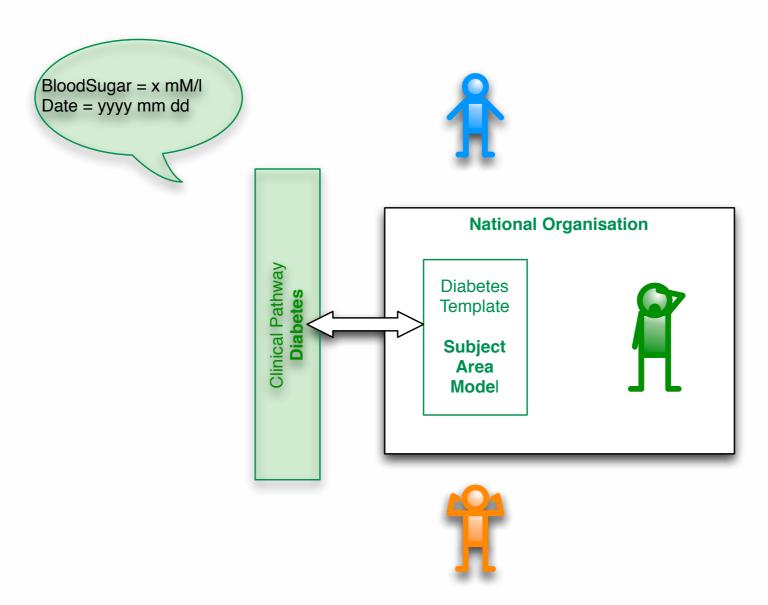


- Standards based eHealth information model which defines
 the representation of the concepts, relationships, constraints, rules, and
 interactions for the HSE's eHealth data.
 It also provides the means to actively control and manage these elements
 across a multitude of national systems.
- When applied. It enables multiple parties to exchange management information about their data sets and their managed data elements.
 It provides an assured, organised, stable and sharable structure of information data for national clinical programmes and records including the enablement of a national EHR.

IA-RM





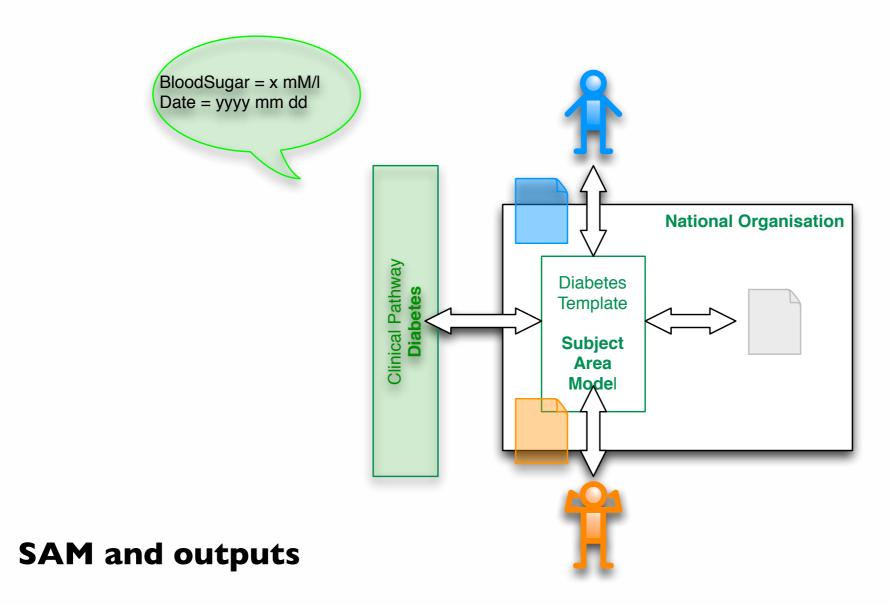


National Organisation

- One National Organisation must be responsible for the IA-RM and its collection of Subject Area Models (SAM's)
- SAM's contain the details that the IT-systems of Actors must adopt



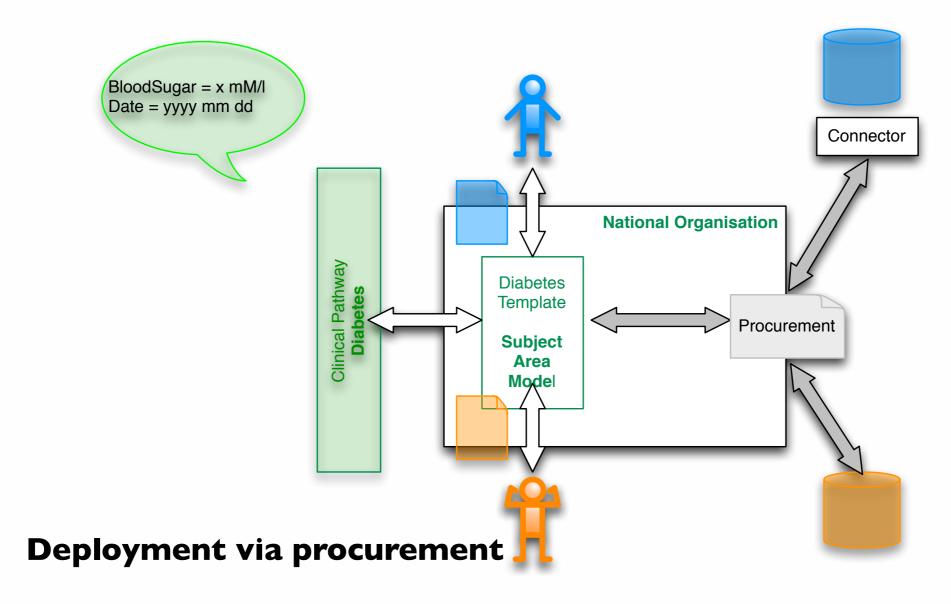




- The SAM's are used to generate outputs that are readable by Humans
- Validation of the SAM's by Actors is essential
- The technical artefacts are generated, also, and help IT-vendors to deploy the SAM's uniformly



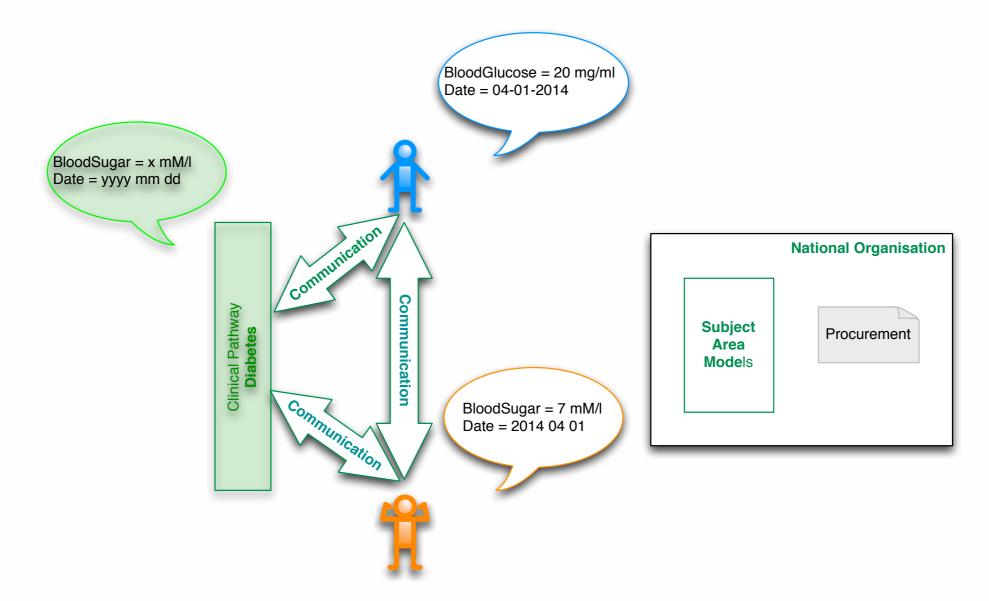




- Procurement is the most efficient way to deploy the IA-RM and SAM's
- Either existing IT-Systems use the SAM's (and outputs)
 to create Connectors
 in order to be able to claim compliance with the standards
- or create new IT-systems that can claim compliance with the standards





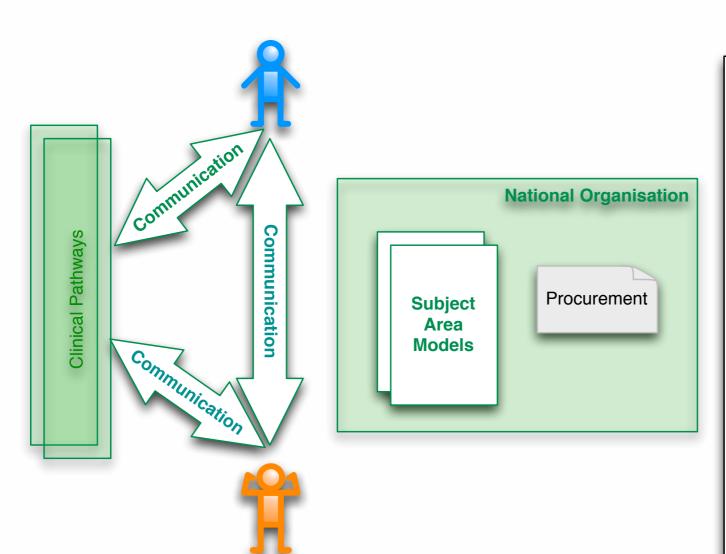


Deployment in healthcare

- Healthcare Actors, using compliant systems, can exchange agreed health data
- Their IT-systems are semantically interoperable







Health Informaticians Knowledge / Experience

- Structures
- Coding systems
- Healthcare

Tooling / services

- SAM editor
- Document Manager
- Integration tools
- Terminology service
- Licenses

Governance

- Relevant Base Standards list
- Production
- Validation of SAM's
- Publication
- Maintenance
- Support
- Validation of compliance testing
- International co-operation

International cross border exchange

- National Contact Point

- The National Organisation has well defined responsibilities
- The new eHealth organisation will need the IA-RM to carry out its functions

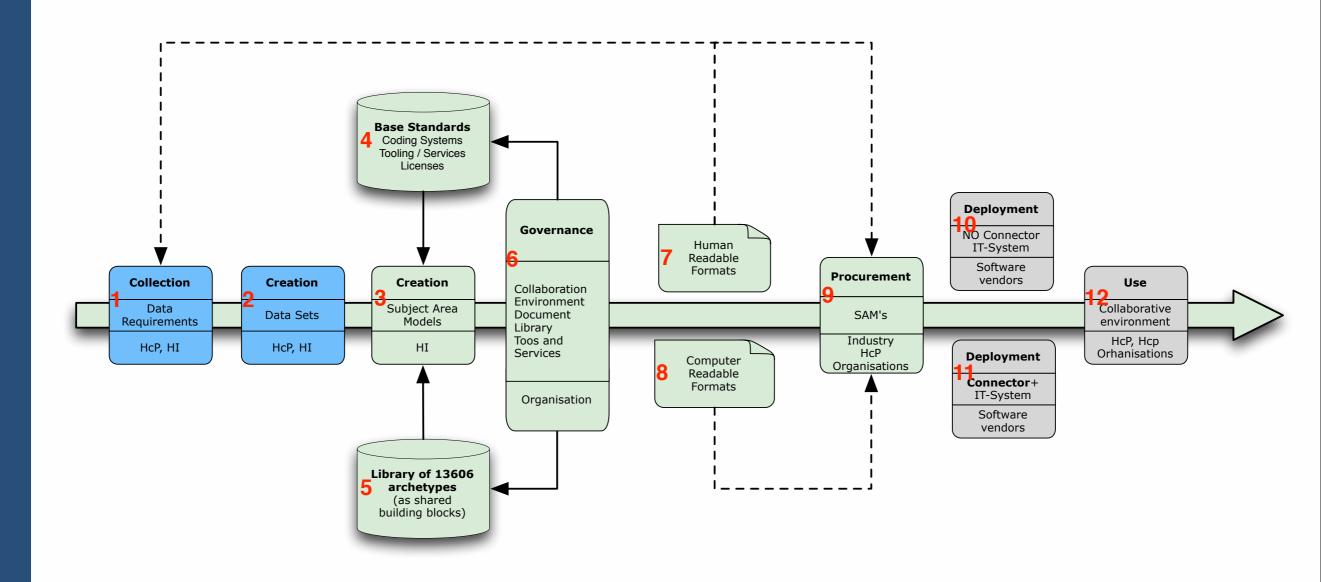


From Clinical Pathways to Collaboration

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Subject Area Models from Clinical Pathways to Collaboration









I - Data requirements

- When Healthcare Providers co-operate they need to agree on the data requirements
- Health Informaticians support Healthcare Providers with the data gathering.
- Healthcare Professional Organisations will play a role





2 - Data Set

- After consensus the Data Set is created based on the acquired data requirements
- Healthcare Providers and their Organisations are facilitated by the Health Informatician





3 - **SAM**

- Health Informaticians define the Subject Area Models (SAM'S)
- These are CEN ISO 13606 Templates,
 built using CEN ISO 13606 Archetypes
- The SAM's express more details than ISO 11179 Data Dictionaries allow

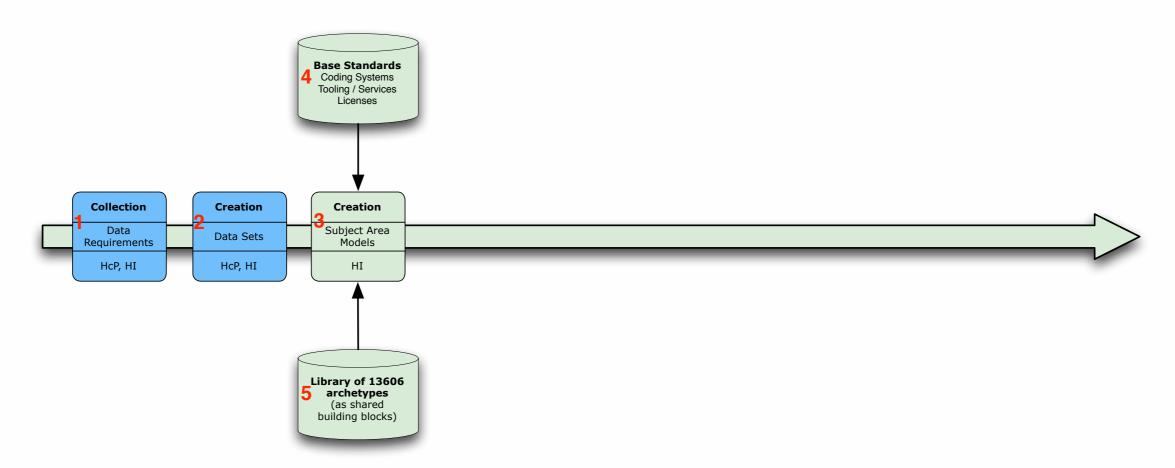




4 - RESOURCES

- SAM's are based on the CEN ISO EHR Communication Standard
- and many supporting relevant adopted standards
- including licensed Coding Systems such as: SNOMED-CT, ICD-x, LOINC, etc.
- Technically the SAM's are CEN ISO 13606 Templates

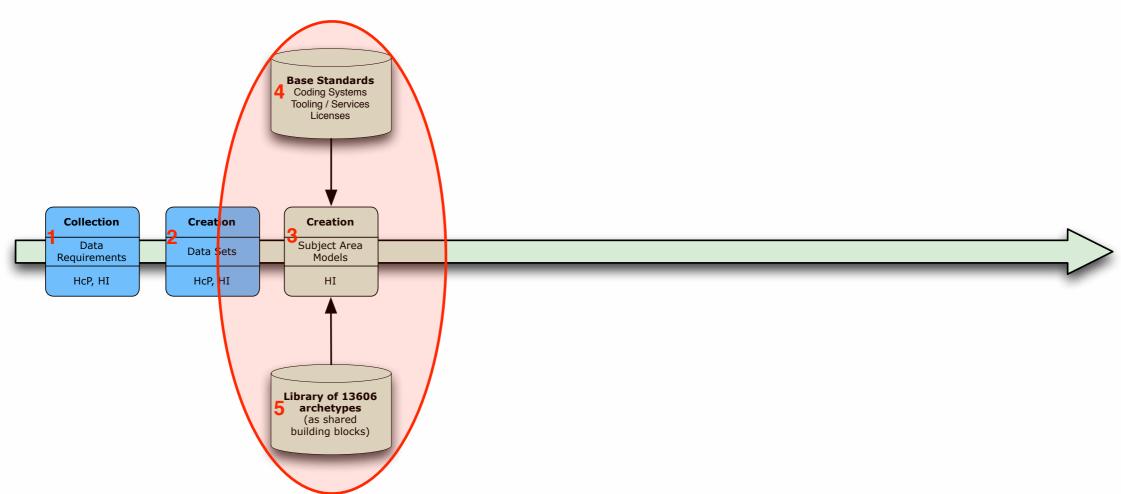




5 - Library of Archetypes as building blocks

- SAM's (13606 Templates) are build using predefined archetypes as shared building blocks
- SAM's will carry relevant codes from the Coding Systems selected
- SAM's represent fully the content of Data Sets



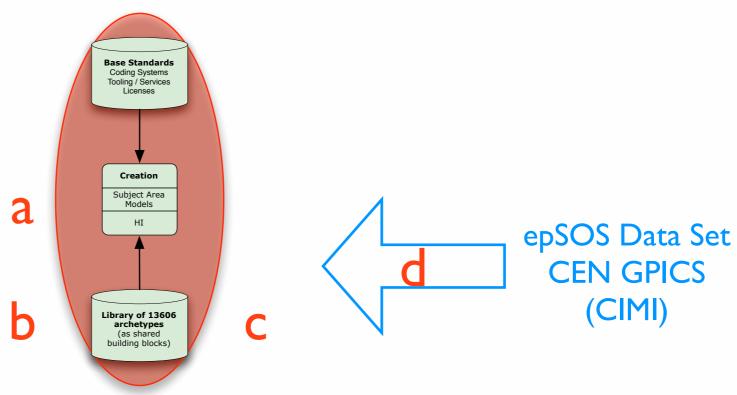


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SAM construction



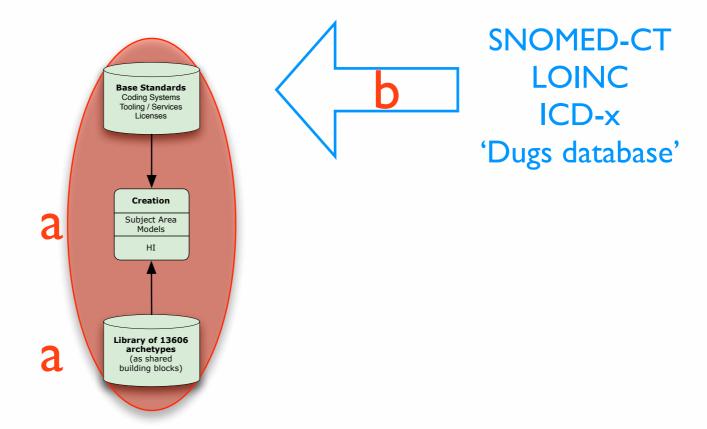


Subject Area Models

- a. Subject Area Models are En I 3606 Templates created with an Archetype Editor
- b. Templates are build using a Library of Archetypes
- c. Library of Archetypes consists of shared patterns/clinical models (e.g. 'Demographics', 'Lab test', 'Diagnosis', 'Complaint', etc.)
- d. Many Data Sets and Archetypes exist already (epSOS data set, GPICS, eventually CIMI project)

SAM construction



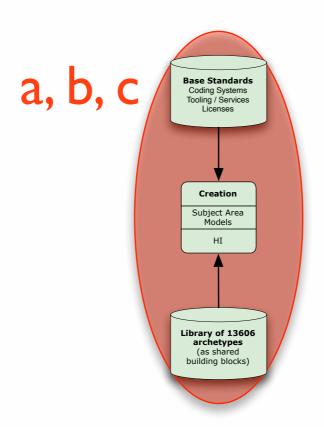


Subject Area Models and Codes

- a. Archetypes and Templates carry codes supplied by a Terminology Server
- b. SNOMED-CT, LOINC, ICD-x, 'Drugs database' (using licenses)

SAM construction

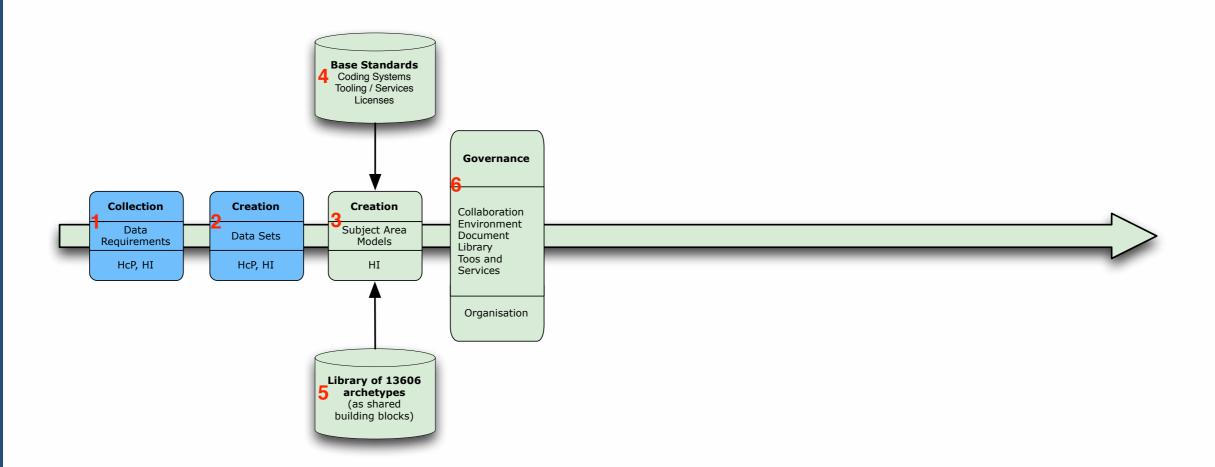




Tooling

- **a. Archetype Editor** to build Templates and Archetypes, and define data elements, attach codes
- **b. Document Management System** to manage documents / files: Archetypes and Templates, to manage the building blocks plus all other associated materials (guides, etc.)
- **c. Terminology Server** to populate the SAM's, Templates, Archetypes with codes

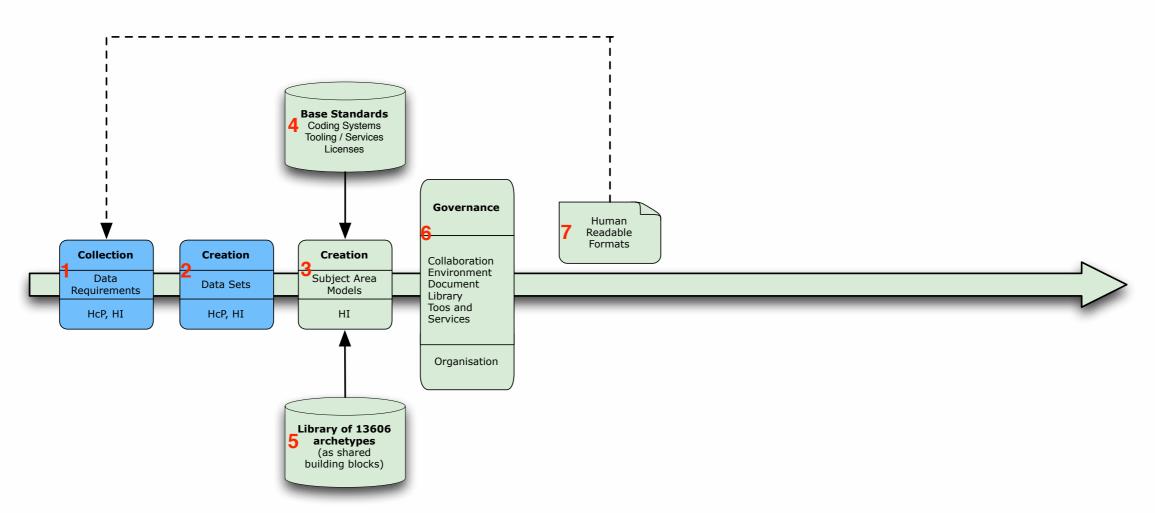




6 - Governance

- An Organisation will govern the process, the tools, services, archetype-, and code set-repositories.
- Observe that one of the services must be a Testing/Validation service to evaluate IT-systems that claim compliance with the published SAM's.

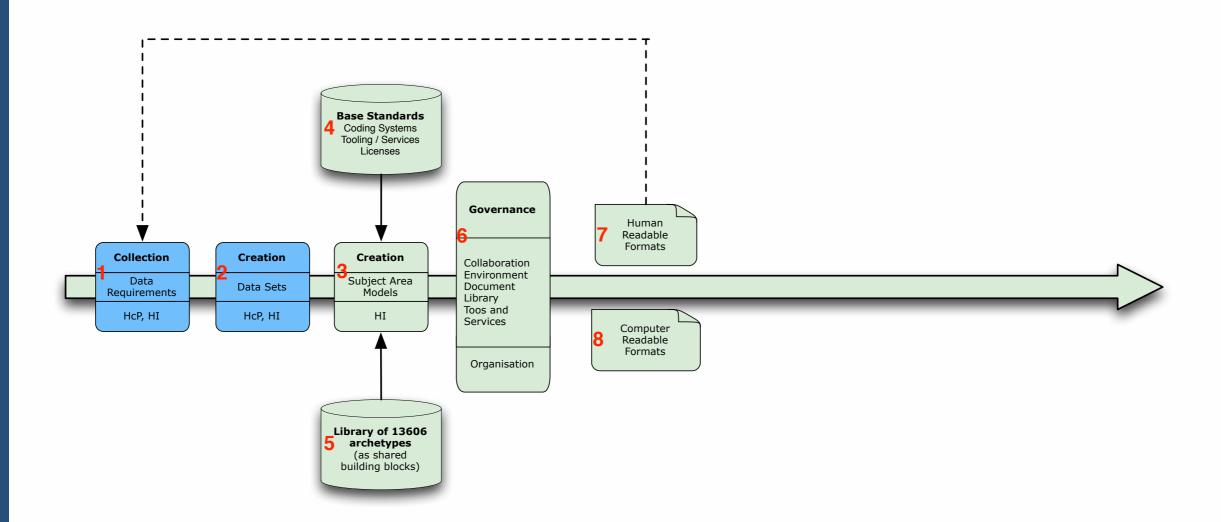




7 - Validation: SAM human readable output

- The SAM editor tool produces human readable output (Spreadsheet, graphs, etc.)
- This output is used by the Health Informatician and the Healthcare Providers and their organisations for the validation of the SAM's produced
- All SAM's an their output will have to be managed in A Document Management System

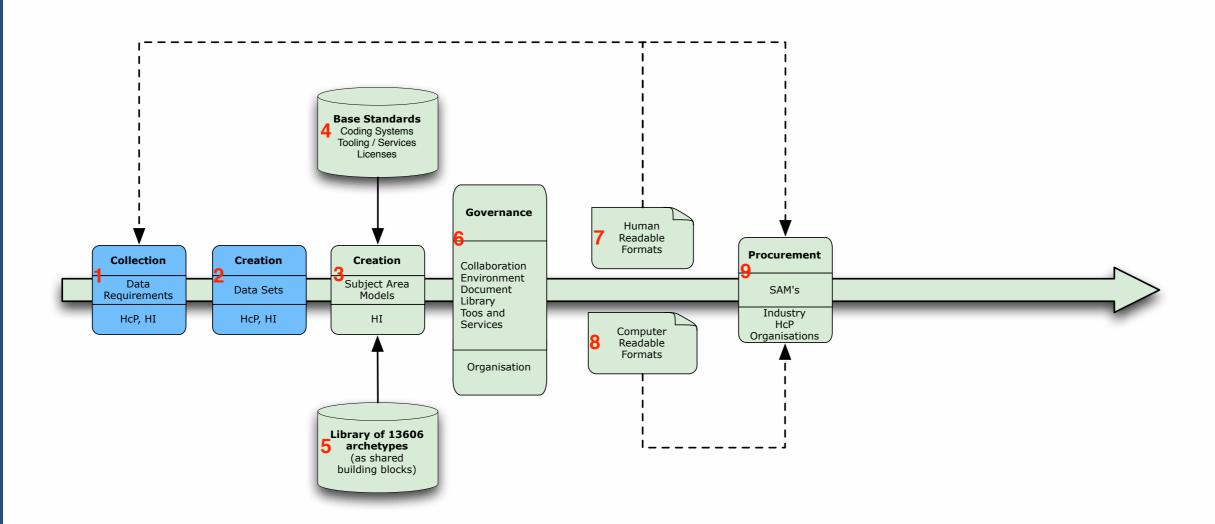




8 - SAM computer procesable output

- Next to human readable output computer processable documents are produced that support the IT-industry with the deployment.
- All SAM's an their output will have to be managed in a Document Management System

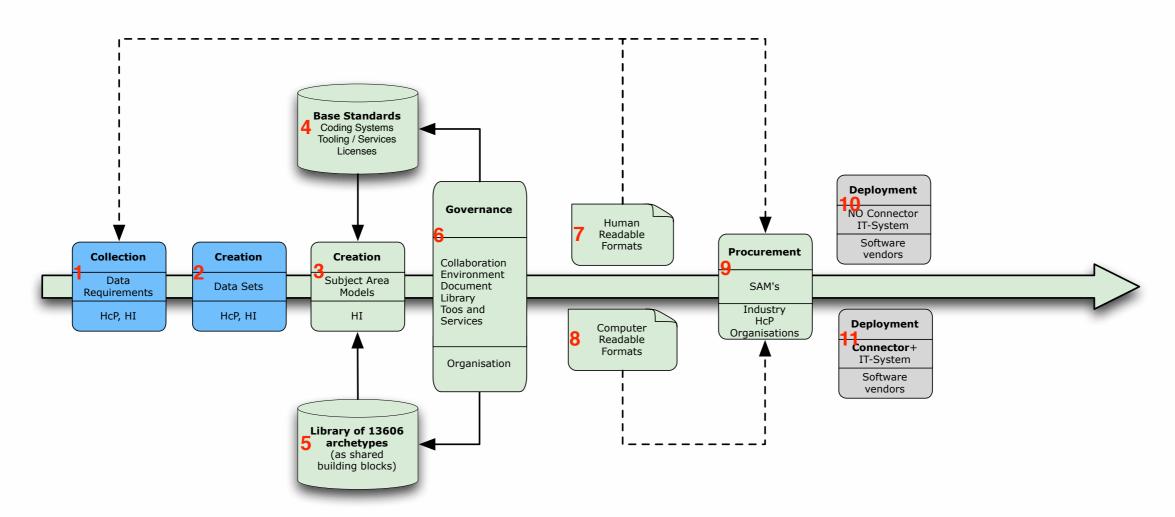




9- Procurement

 All published and validated SAM's and their outputs can play a role in the procurement process for new IT-systems or exchange capabilities.

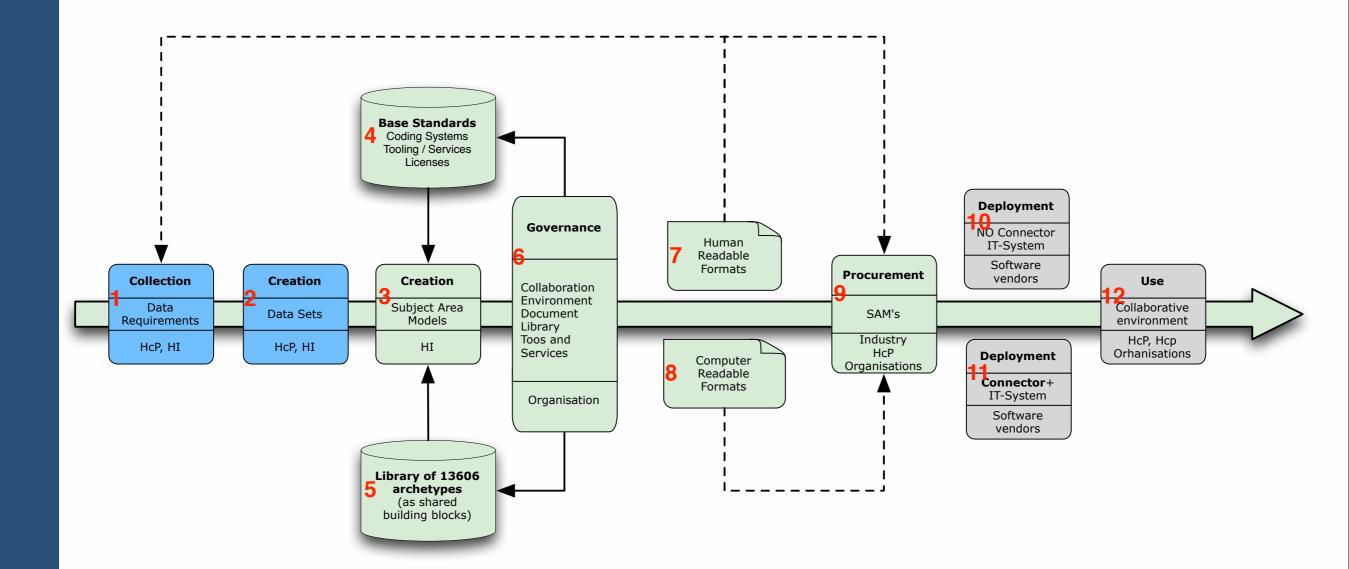




10/11 - Deployment

- The SAM's provide the maximally detailed requirements as expressed by the Healthcare Providers and their organisations based on open International standards
- The IT-industry can use these SAM's to build new systems
- Or exchange solutions (EHR-Extracts, HL7v2, HL7v3 CDA, ...)



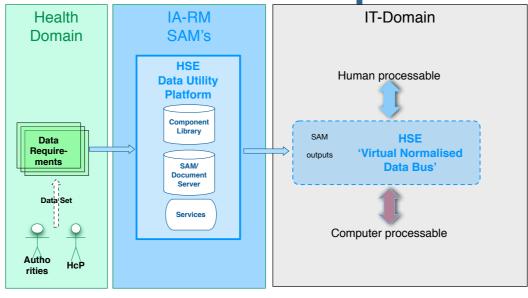


12- Collaborating Healthcare Providers

- IT-systems that deploy the SAM's need to be tested for conformance
- Before these systems are allowed to support collaborating Healthcare Providers

Short term
Next steps



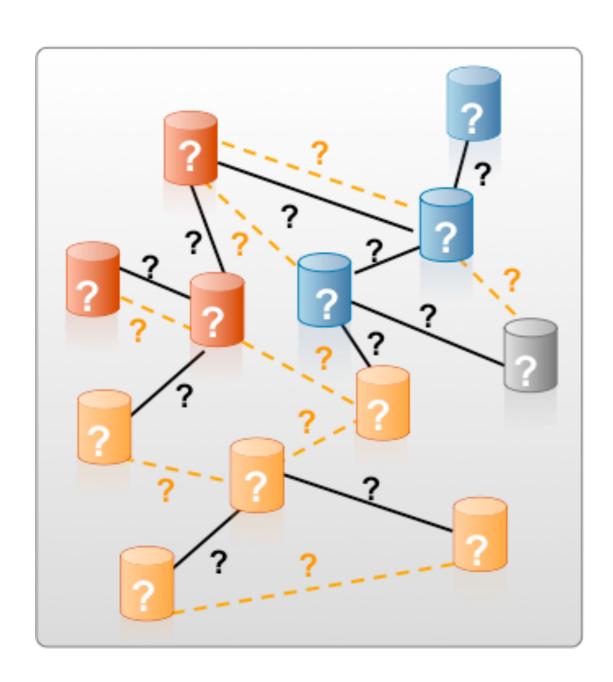


Challenges that can be addressed by IA RM and SAM's and proposed Data Utility Platform:

- uniform Demographics deployment
- e-Prescribing
- Diabetes clinical path way
- MedLis
- Re-Use for clinical research and management reporting
- Data governance and patient safety



Current situation: Big deficits

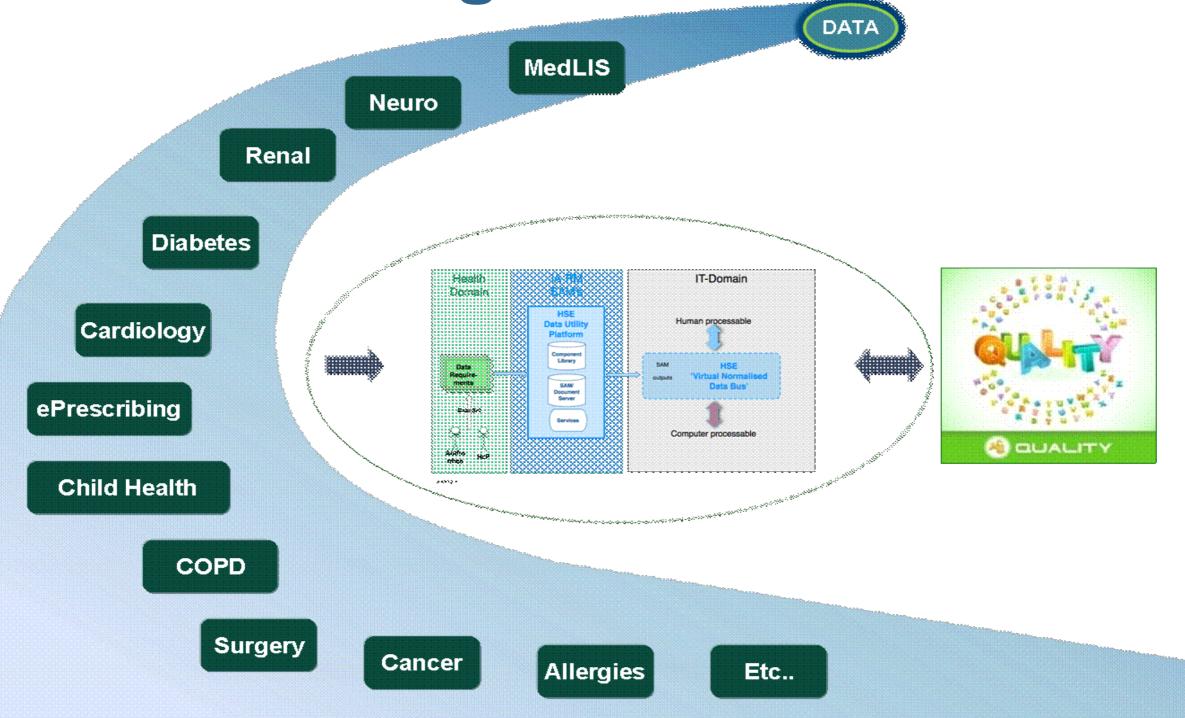


- The Standards based Information Model proposal is based on international best practice
- It provides a means to overcome the current fragmented data landscape of vertical data silos
- As the HSE's information-sharing needs evolve, it provides the means to integrate legacy and new systems whilst supporting accurate identification of multiple entities, including patients, providers and users.

Quality Assured &



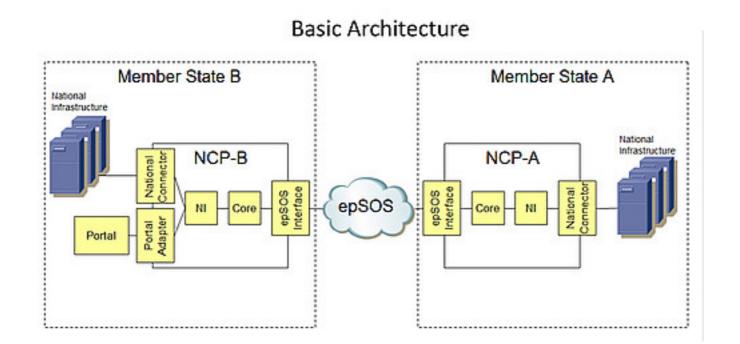




Enables Development of ERS the National Contact Point (NCP) and EHR

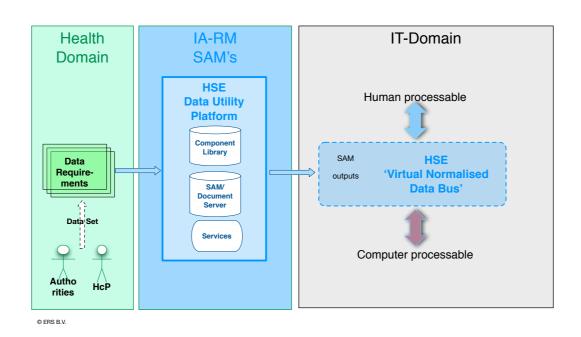


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IA-RM and SAM's Deployment

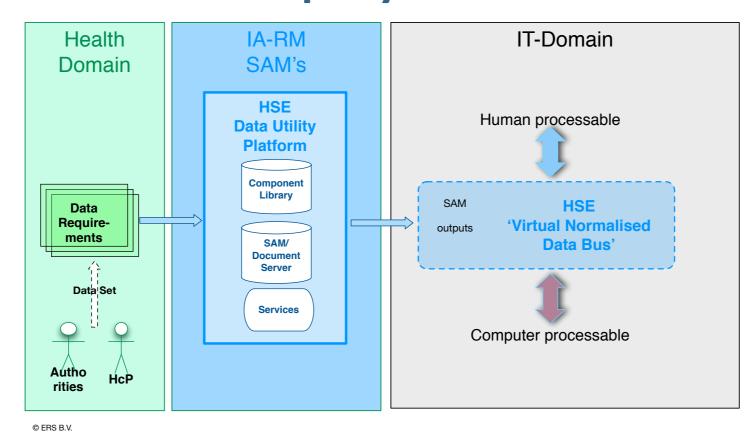


eHealth Organisation needs:

- Tooling
- Professional services



IA-RM and SAM's Deployment



eHealth Organisation cost Possibilities:

- Inside HSE Network: Software licenses, maintenance, professional services
- Inside HSE Network: Outsourced Services



Deployment Next steps

Consider tooling

- SAM editor
- Document Manager
- Terminology Server
- EN13606 conformant database: Clinical Registries
- EN13606 conformant database: Test environment
- Connector(s)
- Professional services