Init1dbw3 - HL7 MDR Requirements (CIC Input)

Pre Interview:

Item	Information/Response
Date:	12/09/2009
Requirement # unique id <*SemCon Ops Initiative*>.<*analysts initials*><*requirement number*> e.g. Init1dbw1 (eventually linked to Use Cases)	Init1dbw3
Originator/Customer's Name:	CBIIT - Margaret Haber
Originator/Customer's Company:	HL7 draft CIC MDR requirements
Stakeholder Community: Enter appropriate category of stakeholder from Primary Stakeholders: • Software and Application designers and architects • Software and Application engineers and developers • Scientific and medical researchers • Medical research protocol designers • Clinical and scientific research data and metadata managers • Clinicians • Patients • Medical research study participants • Broader Stakeholders: caBIG® Community WS NIH projects and related commercial COTS vendors (caEHR, SDO's (HL7, CDISC); International Collaborators (e.g NCRI, cancerGrid, China), Government and regulatory bodies (FDA, CDC, ONC) *(link to view SemConOps Stakeholders description).*	SDO
Summary of requirement pre-interview, by Reviewer:	Support for interoperability between groups with MDRs; Collaboration tools for reviewing similar standards and making refinements to final standard; User friendly Browser; Easy way to create data elements from line of business artifacts such as a spreadsheet or database description; Need to have private and public content in the MDR; Requirements include functional and non-functional (would like to be involved in the requirements process, UI design/feedback sessions for browsers, etc.)
Recommended Next Step Enter one: Follow-up interview, Observe, Use Case Template (text), Use Case Model (formalized/UML diagram), Group Discussion, Prototype, Waiting Room	Review document, schedule interview/questionnaire when Init 1 resources are on board (full time project Business Analysts)