Reaffirm HL7 Version 3 Standard: Implantable Device Cardiac - Follow-up Device Summary, Release 2

Draft

PSS Help and FAQs

Comments

1a. Project Name	Reaffirm HL7 Version 3 Standard: Implantable Device Cardiac - Follow-up Device Summary, Release 2
1b. Project ID	1362
1c. Is Your Project an Investigative Project (aka PSS-Lite)?	No
1d. Is your Project Artifact being Reaffirmed or proceeding to Normative directly after being either Informative or STU?	No
1e. Today's Date	Oct 23, 2019
1f. Name of standard being reaffirmed	HL7 Version 3 Standard: Implantable Device Cardiac - Follow-up Device Summary, Release 2
1g. Project Artifact Information	A project artifact is balloting Normative (no STU) (this includes reaffirmations)
1h. ISO/IEC Standard to Adopt	
1i. Does the standard include excerpted text from one or more ISO, IEC or ISO/IEC standards, but is not an identical or modified adoption?	
1j. Unit of Measure	
2a. Primary/Sponsor WG	Orders & Observations
2b. Co-Sponsor WG	Health Care Devices
2d. Project Facilitator	Lorraine Constable
2e. Other Interested Parties (and roles)	
2f. Modeling Facilitator	
2g. Publishing Facilitator	
2h. Vocabulary Facilitator	

2i. Domain Expert Representative	
2j. Business Requirements Analyst	
2k. Conformance Facilitator	
2l. Other Facilitators	
2m. Implementers	
3a. Project Scope	The therapeutic devices domain comprises the models, messages, and other artifacts that are needed to support messaging related to therapy delivery and observations made by a medical device. Currently, the scope has been focused only on implantable cardiac devices (pacemakers, defibrillators, etc.). This message is related to the follow-up of an Implantable Cardiac Device (pacemaker, defibrillator, etc.) that will contain a subset of device observations, current device therapy settings and device diagnostic information.
Attachments	
3b. Project Need	To enable a standardized device report format to be used across the (Implantable Device Cardiac) IDC sector. Users: IDC industry and cardiologists/cardiac technicians.
3c. Security Risk	
3d. External Drivers	
3e. Objectives/Deliverables and Target Dates	The earliest ballot cycle the TSC will allow - either Feb2020 or May 2020
3f. Common Names / Keywords / Aliases:	
3g. Lineage	
3h. Project Dependencies	
3i. HL7-Managed Project Document Repository URL:	http://www.hl7.org/implement/standards/product_brief.cfm?product_id=41
3j. Backwards Compatibility	Yes
3k. Additional Backwards Compatibility Information (if applicable)	
3l. Using Current V3 Data Types?	Yes
3l. Reason for not using current V3 data types?	
3m. External Vocabularies	Yes
3n. List of Vocabularies	Uses IEEE 11073-10103 standardised IDC terminology
3o. Earliest prior release and/or version to which the compatibility applies	
4a. Products	V2 Messages - Clinical, V3 Messages - Clinical

4b. For FHIR IGs and FHIR Profiles, what product version(s) will the profiles apply to?	
4c. FHIR Profiles Version	
4d. Please define your New Product Definition	
4d. Please define your New Product Family	
5a. Project Intent	Reaffirmation of a standard
5a. White Paper Type	
5a. Is the project adopting/endorsing an externally developed IG?	
5a. Externally developed IG is to be (select one)	
5a. Specify external organization	
5a. Revising Current Standard Info	
5b. Project Ballot Type	Normative (no STU)
5c. Additional Ballot Info	reaffirmation
5d. Joint Copyright	No
5e. I understand I must submit a Joint Copyright Letter of Agreement to the TSC in order for the PSS to receive TSC approval.	no
6a. External Project Collaboration	
6b. Content Already Developed	
6c. Content externally developed?	
6d. List Developers of Externally Developed Content	
6e. Is this a hosted (externally funded) project?	
6f. Stakeholders	Clinical and Public Health Laboratories
6f. Other Stakeholders	
6g. Vendors	Health Care IT
6g. Other Vendors	
6h. Providers	
6h. Other Providers	

6i. Realm	Universal
7d. US Realm Approval Date	
7a. Management Group(s) to Review PSS	
7b. Sponsoring WG Approval Date	Oct 24, 2019
7c. Co-Sponsor Approval Date	
7c. Co-Sponsor 2 Approval Date	
7c. Co-Sponsor 3 Approval Date	
7c. Co-Sponsor 4 Approval Date	
7c. Co-Sponsor 5 Approval Date	
7c. Co-Sponsor 6 Approval Date	
7c. Co-Sponsor 7 Approval Date	
7c. Co-Sponsor 8 Approval Date	
7c. Co-Sponsor 9 Approval Date	
7c. Co-Sponsor 10 Approval Date	
7e. CDA MG Approval Date	
7f. FMG Approval Date	
7g. V2 MG Approval Date	
7h. Architecture Review Board Approval Date	
7i. Steering Division Approval Date	Nov 07, 2019
7j. TSC Approval Date	