ISO IDMP Task Force: International activities and IDMP

EU ISO IDMP Task Force meeting
25 September 2015

Context & Background

ISO IDMP Standards	• Define the required data elements and their structure • Provides 'business-level' description of IDMP	ISO TC/215 WG6
ISO IDMP Implementation Guides (Technical Standards)	• Define the technical details on how to implement the standards • Includes field formats, business rules etc.	ISO TC/215 WG6
HL7 Messaging Specifications	• Defines the messages that will be used to exchange IDMP information • Based on existing HL7 *'Common Product Model'* standard (similar to FDA's SPL)	HL7 WG
EU Implementation Guide	• Interpretation of fields specifically for EU regulatory environment • Guidance on processes of submitting and updating data	EMA (P&S Project)

High level Project Plan (update in progress)

	2015			2016								
	Oct	Nov	Dec	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep
ISO IDMP Standards												
11238 Substance *(Update to original)*			Draft							Publication		
11239 Dose, pres. units, routes, packaging	*Final and stable*											
11240 Units of Measure	*Final and stable*											
11615 Medicinal Product *(Update)*			Draft							Publication		
11616 Pharmaceutical Product *(Update)*			Draft							Publication		
ISO IDMP Implementation Guides (Technical Standards)												
11238 Substance (Core + 4 annexes)	*Final and stable*											
11238 Substance (Remaining 6 annexes)		Draft								Publication		
11239 Dose, pres. units, routes, packaging		Draft			Publication							
11615 Medicinal Product		Draft			Publication							
11616 Pharmaceutical Product		Draft			Publication							
HL7 Messaging Formats												
Substance	Publication											
Product	Publication											