

Initial Gap Analysis on Standards Development

Presentation for HL7 International Work Group Biomedical Research & Regulation (BR&R)
Tuesday 22 September 2020 – Q2



Work Package 1 - IDMP related standards and terminologies

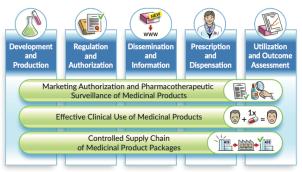


► Making IDMP and related standards and terminologies work across the landscape of implementation

3 **Application** fields



Combining the perspectives



Implementation domains











Dissemination

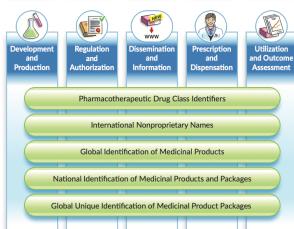
and







Utilization





Initial Gap Analysis – through workshops, wish lists, and use cases from UNICOM participants



Use cases:

- ▶ ePrescription
- □ Adverse Events (reporting to the authorities and beyond)
- ▶ Medication Errors

► Audience:

Standards Developing Organizations

Process:

- ▶ WP-1 to coordinate gap analysis across use cases and implementation domains
- ▶ WP-1 to hand over to the relevant SDO's to address an interim and final solution
- SDO's to discuss the identified gaps, with help from WP-1 and other informed experts
- > SDO's to formulate a response, including estimated timeline for final resolution
- > WP-1 to disseminate this response an possible interim solutions within UNICOM
- ▶ WP-1 to inform and discuss with the Community of Expertise where appropriate























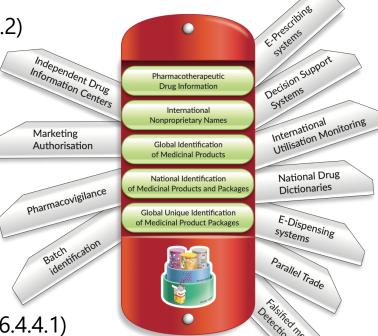


Identified gaps across implementation domains – proposed next steps within HL7 International



- ► Evaluate impact and implement new manufactured item class / identifier (6.2.3.2)
- ▶ IDMP & FHIR alignment cross-workgroup use cases and requirements definition (6.2.7.1)
- ► IDMP & FHIR alignment resource harmonisation (6.2.7.3)
- ▶ Development of dispensation feedback message to prescriber including DCID (6.3.1.2)
- ► Logical model requirements from different use cases (6.3.6.x)
- ▶ Profile/IG updates for ePrescribing and eDispensing to align to IDMP (6.3.7.2)
- ▶ eDispensing related items and attributes in IDMP (6.4.1.1)
- ► Adverse Events processes, structured data and value sets (6.4.2.2)
- ► Medication Error processes, structured data and value sets (6.4.2.4)
- ► Guidance on the use of IDMP on mHealth apps (6.4.3.1)

- ▶ Bridging the regulatory and clinical domains quality and maturity considerations (6.4.4.1)
- ▶ Use of IDMP to improve trust in AI through structured data (6.4.4.2)







Let's work together to make this happen!

