ISO/TR 14872:2025-12 (E)

Health informatics - Identification of medicinal products - Core principles for maintenance of identifiers and terms

Contents

Page

Foreword iv

Introduction v

1 Scope 1

2 Normative references 1

3 Terms and definitions 1

4 Abbreviated terms 2

5 Maintenance of International Standards on IDMP -- description of existing relationships between regulators and other standards development organizations 2

6 IDMP terminology maintenance 3
6.1 IDMP maintenance organizations at global level 3
6.2 Core data within IDMP and data that is "derived" 3

7 Current maintenance processes 3
7.1 Generic description of governance and maintenance process 3
7.2 Governance and maintenance: substances and Global Pharmaceutical Product Identifier (PhPID) 4
7.3 Governance and maintenance: pharmaceutical dose forms 5
7.4 Governance and maintenance: units of measurement 5
7.5 Governance for SNOMED interactions with IDMP 6
7.6 Governance for MedDRA 7
7.7 Governance for Anatomical Therapeutic Chemical (ATC) classification system and Defined Daily Dose (DDD) 8
7.8 Governance for International Nonproprietary Names (INN) 9
7.9 Governance for GS1 10

8 IDMP core maintenance processes 11
8.1 General 11
8.2 General processes 11
8.3 Terminology processes 11
8.4 Unambiguity 12

9 International mapping and language translations 12

10 Integration of IDMP core with data sources that are considered "derived" 12

Bibliography 14