



Coding with MedDRA



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MedDRA was developed under the auspices of the International Council for Harmonisation of Technical Requirements for Pharmaceuticals for Human Use (ICH). The activities of the MedDRA Maintenance and Support Services Organization (MSSO) are overseen by an ICH MedDRA Management Committee, which is composed of the ICH parties, the Medicines and Healthcare products Regulatory Agency (MHRA) of the UK, Health Canada, and the WHO (as Observer).

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Course Overview

- MedDRA background
- MedDRA's structure, scope, and characteristics
- MedDRA maintenance
- Coding conventions
- Synonym lists
- Quality assurance (QA) of coding
- MedDRA Term Selection: Points to Consider document
- Hands-on coding exercises

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MedDRA Background



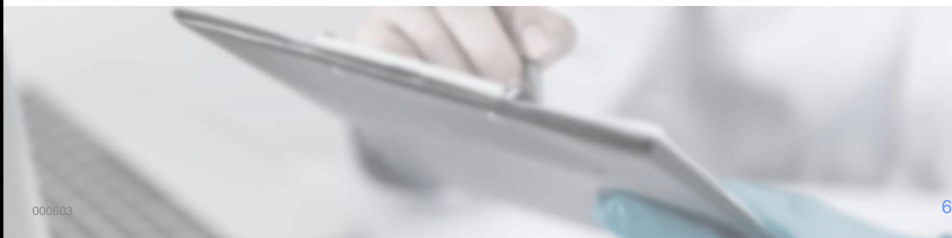
What is MedDRA?

Med = Medical

D = Dictionary for

R = Regulatory

A = Activities





MedDRA

MedDRA Definition

MedDRA is a clinically-validated international medical terminology used by regulatory authorities and the regulated biopharmaceutical industry. The terminology is used through the entire regulatory process, from pre-marketing to post-marketing, and for data entry, retrieval, evaluation, and presentation.

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MedDRA

MedDRA's Purpose

- Facilitate the exchange of clinical information through standardization
- Important tool for product evaluation, monitoring, communication, electronic records exchange, and oversight
- Supports coding (data entry) and retrieval and analysis of clinical information about human medical products including pharmaceuticals, biologics, vaccines, and drug-device combination products

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MedDRA

MedDRA and the MSSO

- International support and development of terminology
- Foster use of MedDRA through communications and educational offerings
- “Custodians”, not owners, of the terminology
- JMO (partner organization for Japanese-language MedDRA)
- Governed by a Management Committee (industry, regulators, multi-national, other interested parties)

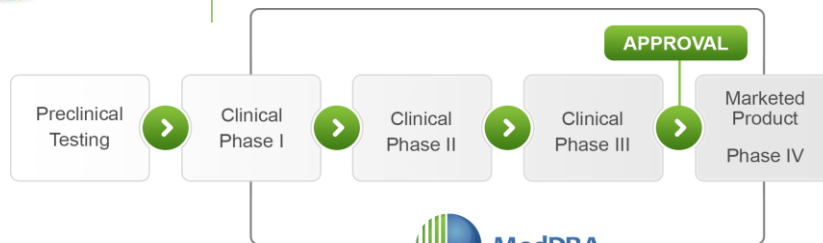
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MedDRA

Where MedDRA is Used



Regulatory Authority and Industry Databases
Individual Case Safety Reports and Safety Summaries

Clinical Study Reports

Investigators' Brochures

Core Company Safety Information

Marketing Applications

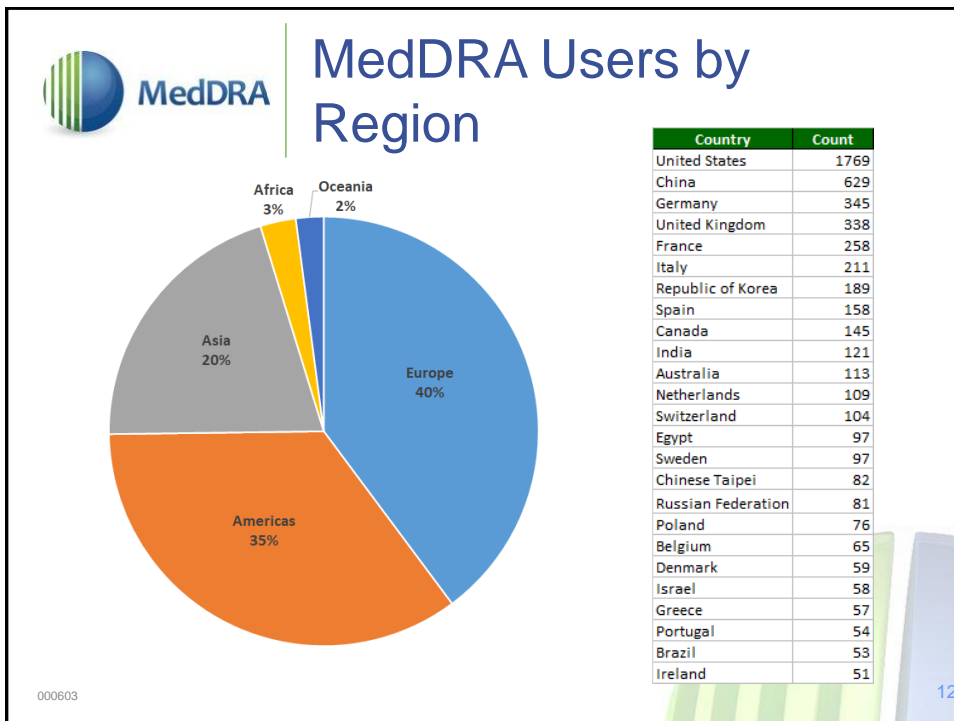
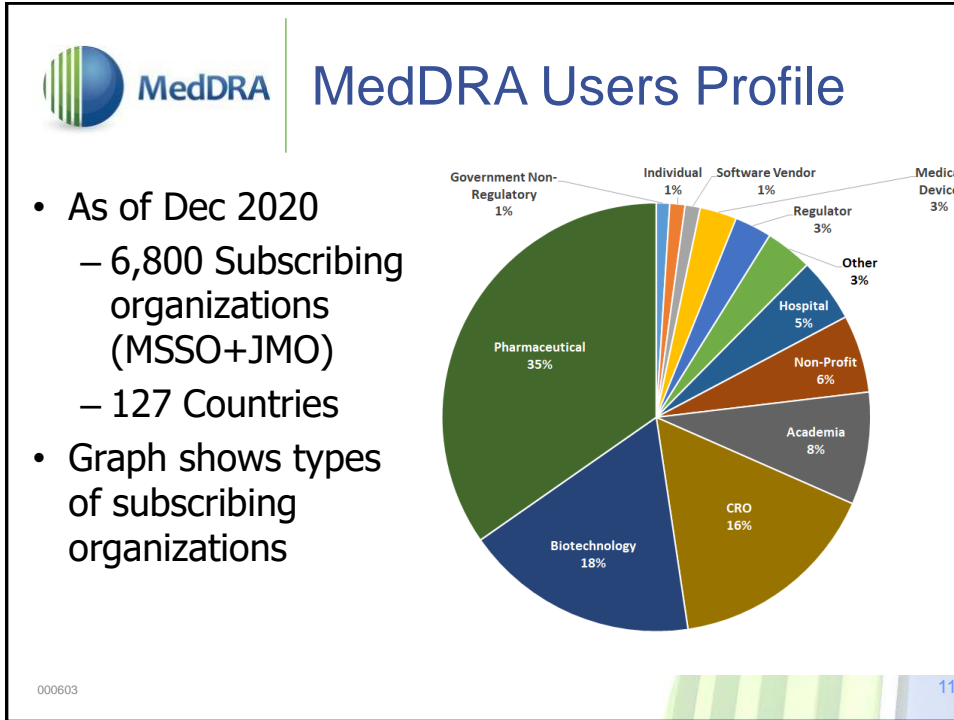
Publications


Prescribing Information

Advertising


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


MedDRA Data Sharing



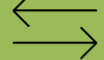
Access

Subscription grants access to MedDRA for one year




3rd Party Distribution

Subscriber cannot grant any sublicense, publish or otherwise distribute MedDRA to a third party



Free Exchange


Data may be freely exchanged between current MedDRA subscribers



Violation

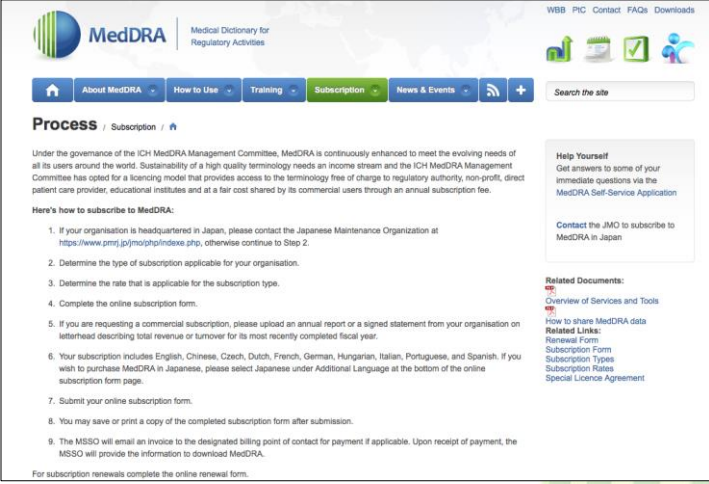
Sharing MedDRA with a non-subscribing organization is a violation of the MedDRA license

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MedDRA Data Sharing (cont)

- For details, see the Statement on MedDRA Data Sharing



Process / Subscription / #

Under the governance of the ICH MedDRA Management Committee, MedDRA is continuously enhanced to meet the evolving needs of all its users around the world. Sustainability of a high quality terminology needs an income stream and the ICH MedDRA Management Committee has opted for a licensing model that provides access to the terminology free of charge to regulatory authority, non-profit, direct patient care provider, educational institutes and at a fair cost shared by its commercial users through an annual subscription fee.

Here's how to subscribe to MedDRA:

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7. Submit your online subscription form.
8. You may save or print a copy of the completed subscription form after submission.
9. The MSSO will email an invoice to the designated billing point of contact for payment if applicable. Upon receipt of payment, the MSSO will provide the information to download MedDRA.

For subscription renewals complete the online renewal form.

Help Yourself
Get answers to some of your immediate questions via the MedDRA Self-Service Application

Contact the JMO to subscribe to MedDRA in Japan

Related Documents:


- Overview of Services and Tools
- How to share MedDRA data
- Renewal Form
- Subscription Form
- Subscription Types
- Subscription Rates
- Special License Agreement

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MedDRA's Scope, Structure, and Characteristics

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Scope of MedDRA

Not a drug dictionary

Patient demographic terms

Clinical trial study design terms

OUT

IN

Medical conditions

Indications

Investigations (tests, results)

Medical and surgical procedures

Medical, social, family history

Medication errors

Product quality issues

Device-related issues

Product use issues

Pharmacogenetic terms

Toxicologic issues

Standardized queries

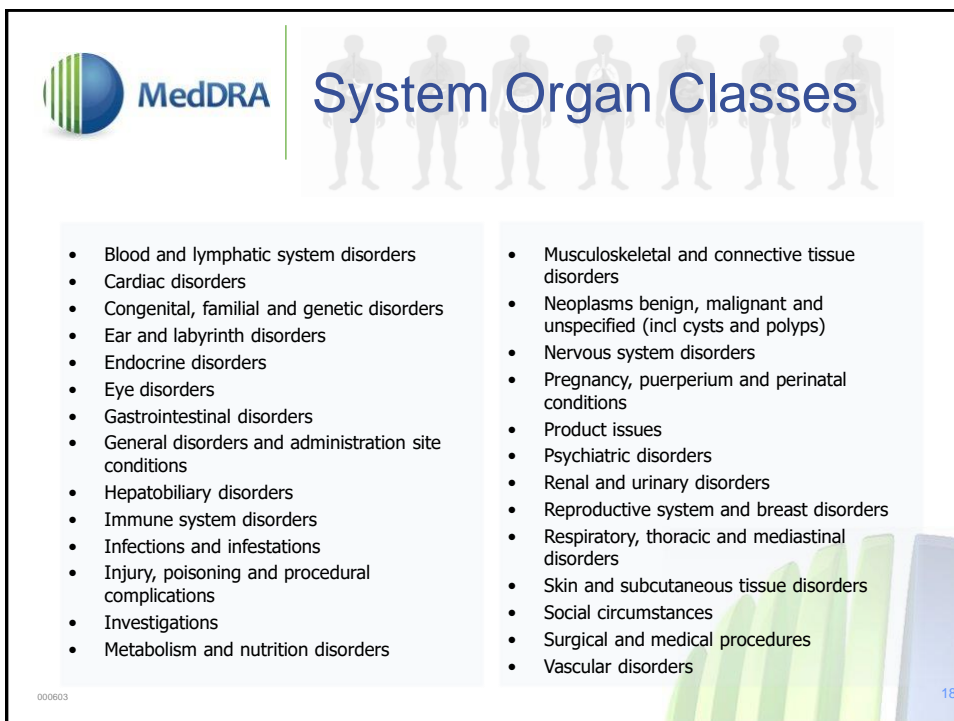
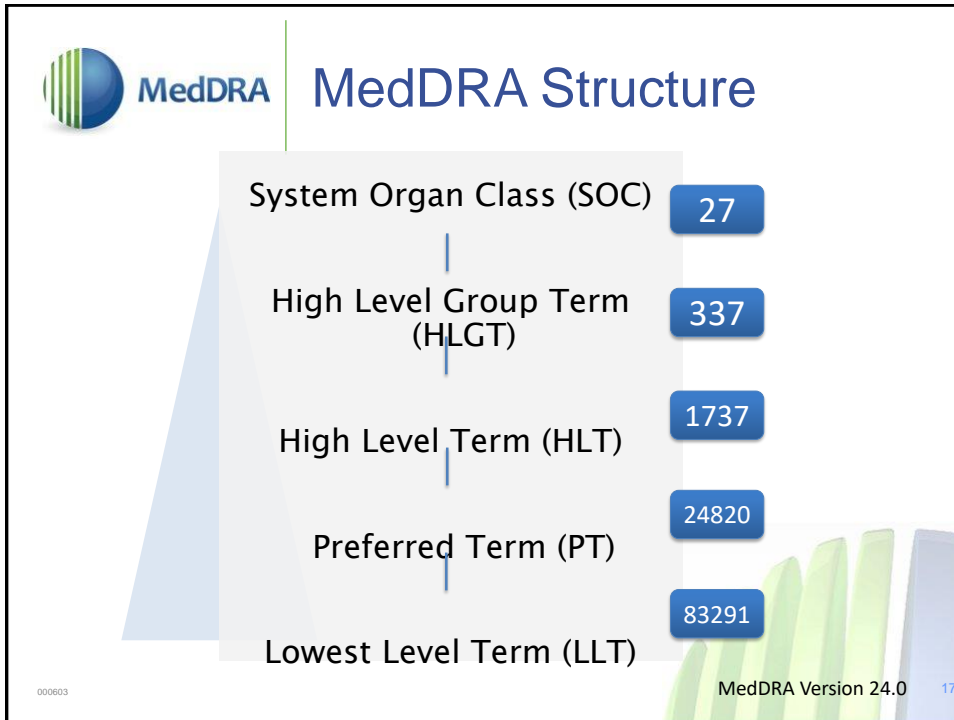
Frequency qualifiers

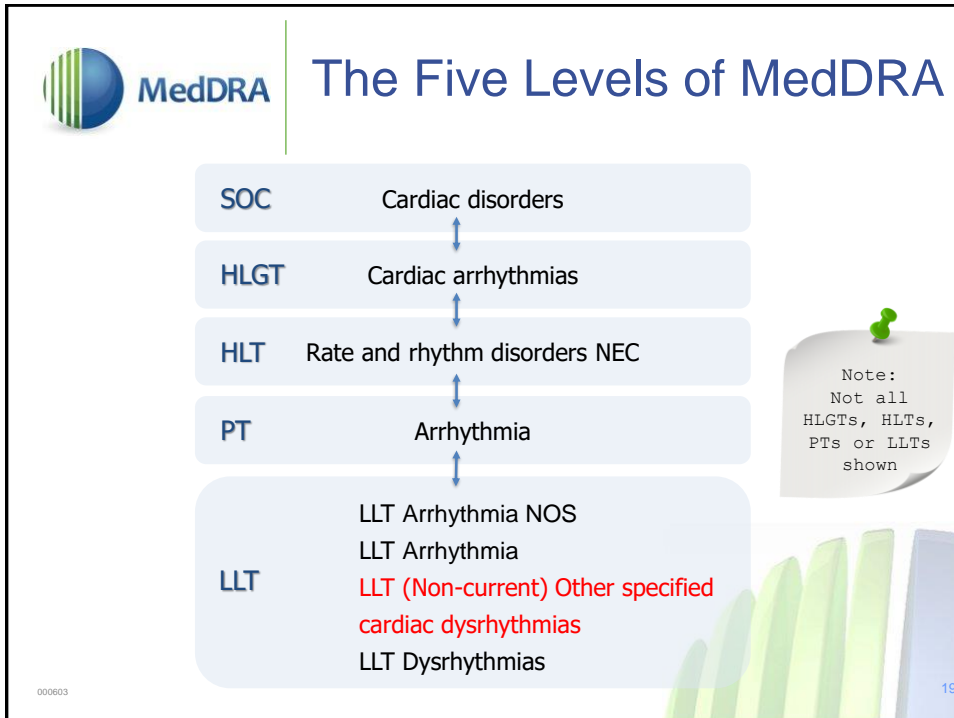
Numerical values for results

Severity descriptors

Not an equipment, device, diagnostic product dictionary

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MedDRA Non-Current Terms LLT

- Flagged at the LLT level in MedDRA
- Not recommended for continued use
- Retained to preserve historical data for retrieval and analysis
- Terms that are vague, ambiguous, outdated, truncated, or misspelled
- Terms derived from other terminologies that do not fit MedDRA rules


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MedDRA Codes


- Each MedDRA term assigned an 8-digit numeric code starting with "1"
- The code is non-expressive
- Codes can fulfill a data field in various electronic submission types (e.g., E2B)
- New terms are assigned sequentially

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Codes and Languages

Kopfschmerz **German** Fejfájás **Hungarian**
 Cefaleia **Portuguese** Cefalea **Italian**
 Hoofdpijn **Dutch** 頭痛 **Japanese**
 Headache **English** Cefalea **Spanish**
 Céphalée **French** Головная боль **Russian**
 Bolest hlavy **Czech** 두통 **Korean**
 头痛 **Chinese** Cefaleia **Brazilian Portuguese**

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 Electronic Submission

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MedDRA

A Multi-Axial Terminology

Multi-axial = the representation of a medical concept in multiple SOC

- ✓ Allows grouping by different classifications
- ✓ Allows retrieval and presentation via different data sets

All PTs assigned a primary SOC

- ✓ Determines which SOC will represent a PT during cumulative data outputs
- ✓ Prevents "double counting"
- ✓ Supports standardized data presentation
- ✓ Pre-defined allocations should not be changed by users

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MedDRA

A Multi-Axial Terminology (cont)

SOC = Respiratory, thoracic and mediastinal disorders
(Secondary SOC)

SOC = Infections and infestations
(Primary SOC)

HLGT = Respiratory tract infections

HLGT = Viral infectious disorders


HLT = Viral upper respiratory tract infections

HLT = Influenza viral infections

PT = Influenza

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Rules for Primary SOC Allocation




- PTs represented in only one SOC are automatically assigned that SOC as primary
- PTs for diseases, signs and symptoms are assigned to prime manifestation site SOC
- Congenital and hereditary anomalies terms have SOC *Congenital, familial and genetic disorders* as Primary SOC
- Neoplasms terms have SOC *Neoplasms benign, malignant and unspecified (incl cysts and polyps)* as Primary SOC
 - **Exception:** Cysts and polyps have prime manifestation site SOC as Primary SOC
- Infections and infestations terms have SOC *Infections and infestations* as Primary SOC

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Primary SOC Priority

If a PT links to more than one of the exceptions, the following priority will be used to determine primary SOC:

-  *Congenital, familial and genetic disorders*
-  *Neoplasms benign, malignant and unspecified (incl cysts and polyps)*
-  *Infections and infestations*

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MedDRA

A Multi-Axial Terminology (cont)

PTs in the following SOC **only** appear in that particular SOC and not in others, i.e., they are not multi-axial

- *Investigations*
- *Surgical and medical procedures*
- *Social circumstances*

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MedDRA

Can You Select the Primary SOC for This PT?

PT	HLT	HLGT	SOC
Congenital HIV infection	Viral infections congenital	Infections and infestations congenital	Congenital, familial and genetic disorders
	Congenital neonatal infections	Neonatal and perinatal conditions	Pregnancy, puerperium and perinatal conditions
	Retroviral infections	Viral infectious disorders	Infections and infestations
	Acquired immunodeficiency syndromes	Immunodeficiency syndromes	Immune system disorders

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MedDRA Maintenance



MedDRA Maintenance

- Users can send change requests (CRs) to MSSO for consideration
 - Organizations allowed 100 CRs/month
 - Rigorous medical review by MSSO physicians
 - For simple changes (PT and LLT levels), response within 7-10 working days
 - Complex changes (above PT level) posted for comments mid-year
- Two MedDRA updates/year
 - 1 March X.0 (Complex release)
 - 1 September X.1 (Simple release)



MedDRA WebCR

- Web-based tool for Change Requests (CR)
 - URL: <https://mssotools.com/webcr/>
 - Via the Change Request Information page
- Ability to submit CRs online
 - Requests must be in English
- Immediate confirmation
- Review unsubmitted CRs online
- Ability to query CR history back to v5.1

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MedDRA Submitting Changes



- Online change request submission tool guides user to enter all needed information
- Instructional video on use of WebCR on “Training” page of MedDRA website


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MedDRA Submitting Changes (cont)

Add a New PT

Proposed PT (Required) 

Second degree chemical burns of skin

Primary HLT (Optional)

Chemical injuries

Primary SOC (Optional)

Injury, poisoning and procedural complications


Secondary HLT (Optional)

Dermatitis ascribed to specific agent

Secondary SOC (Optional)

Skin and subcutaneous tissue disorders

Justification statement is required

Justification 


Please consider including the gradation of chemical burns similar to the gradation of thermal burns under HLT Thermal burns to assist with coding and analysis.

Attach supporting document (Optional)

Attachment
000603 C:\Users\is804733\Desktop\SupportingInforma Browse...

- Sample entry for a new PT in WebCR
- Justification and supporting documentation is important to help MSSO understand the need

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MedDRA Proactive MedDRA Maintenance

- What is the proactive approach?
 - Corrections/improvements made internally by the MSSO
 - General changes suggested by users
- Submitting ideas
 - Send to MSSO Help Desk. Justification is helpful.
 - Example: Review placement of bruise and contusion terms to facilitate coding and analysis
- Evaluation of proposals
 - Final disposition is not time limited; MSSO may take time to review
 - Proactive approach does not replace usual CR process

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MedDRA

MedDRA Version Analysis Tool (MVAT)

- Web-based (<https://tools.meddra.org/mvat>)
- Free to all users
- Features
 - *Version Report Generator* (produces exportable report comparing any two versions)
 - *Data Impact Report* (identifies changes to a specific set of MedDRA terms or codes uploaded to MVAT)
 - *Search Term Change* (identifies changes to a single MedDRA term or code)

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MedDRA


MedDRA Version Analysis Tool (MVAT) (cont)

- User interface and report output available in all MedDRA languages
- Ability to run reports on supplemental changes
- Option to run reports on secondary SOC changes
- Watch the two-part webinar on the use of MVAT under "Tools" tab on the "Training Materials" page of MedDRA website

<https://www.meddra.org/training-materials>

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MedDRA Version Analysis Tool (MVAT) (cont)

MedDRA Version Analysis Tool (MVAT)

Version Report Description

Select Different Versions to Compare

Language: English

Starting Version: MedDRA 23.1 English

Ending Version: MedDRA 24.0 English

Include Secondary SOC Information

Select SOCs to filter (default is all SOCs)

- Blood and lymphatic system disorders
- Cardiac disorders
- Congenital, familial and genetic disorders
- Ear and labyrinth disorders
- Endocrine disorders
- Eye disorders
- Gastrointestinal disorders
- General disorders and administration site conditions
- Hepatobiliary disorders
- Immune system disorders

Note: The starting MedDRA version must be older than the ending MedDRA version


Preferred Language: English

Release Supplemental View: Release


-
-
-
-

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MSSO's MedDRA Browsers




MedDRA Desktop Browser (MDB)
Download MDB and release files from MedDRA website



Mobile MedDRA Browser
<https://mmb.meddra.org>



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MSSO's MedDRA Browsers (cont)

- Features
 - Each require MedDRA ID and password
 - View/search MedDRA and SMQs
 - Support for all MedDRA languages
 - Language specific interface
 - Ability to export search results and Research Bin to local file system (MDB and WBB only)

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MDB and WBB Special Features

- Preview upcoming (supplemental) changes in next release*
- View primary **and** secondary link information
- Upload terms to run against SMQs
- Advanced search options (e.g., NOT, OR)

*Supplemental view not available on MDB

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Data WBB Supplemental View

- Where will Change Requests I submitted appear in next release of MedDRA?
- What are the changes in an area of MedDRA in the next release?

Change View to Supplementals

The screenshot shows the MedDRA Browser interface. At the top, there is a navigation bar with 'Release/Supplemental View' and 'Release' buttons. A dropdown menu is open, showing 'Supplementals' as the selected option. Below the navigation bar, there are search options and a list of search results.

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WBB Supplemental View (cont)

- Display changes color to pink
- Supplemental terms highlighted by broken green line
- Changes for the next release are displayed immediately after they are approved by the MSSO

The screenshot shows the MedDRA Browser - Supplemental View interface. The header is pink. The search results are displayed in a table with columns for MedDRA Code, MedDRA Term, and Category. The table shows results for 'Device delivery system' terms.

MedDRA Code	MedDRA Term	Category
10077122	Device delivery system issue	Y

Below the table, there is a section for 'LLT Occurrences in MedDRA (Supplemental)' with a list of terms and their categories.

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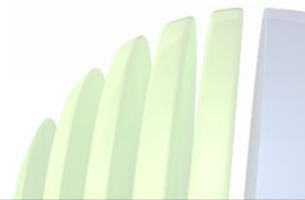
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MedDRA Browser Demonstration and Instruction



Coding Exercises

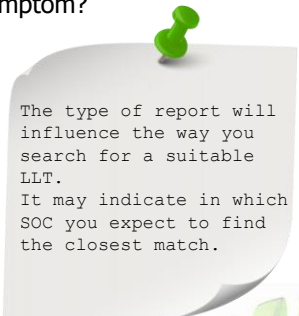




MedDRA

Assessing the Reported Information

- Consider what is being reported. Is it a:
 - Clinical condition - Diagnosis, sign or symptom?
 - Indication?
 - Test result?
 - Injury?
 - Procedure?
 - Medication error?
 - Product use issue?
 - Product quality issue?
 - Social circumstance?
 - Device issue?
 - Procedural complication?
- **Is it a combination of these?**



The type of report will influence the way you search for a suitable LLT. It may indicate in which SOC you expect to find the closest match.

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MedDRA

MedDRA Browsing Tips

- First, try using actual words from reporter
- Use “top-down” and “bottom-up” approaches
- Look at the “neighbors” and check the hierarchy
- Consider synonyms, e.g., “Liver” and “Hepatic”
- Use word stems, e.g., “Pancrea”

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MedDRA

MedDRA Browsing Tips (cont)

- Use available resources for difficult verbatim terms (web search, medical dictionaries, colleagues)
- Use advanced Boolean search terms features (i.e., "begins with", "exact match", "ends with", "not contains", "and", "or") when needed
- Become familiar with MedDRA Concept Descriptions

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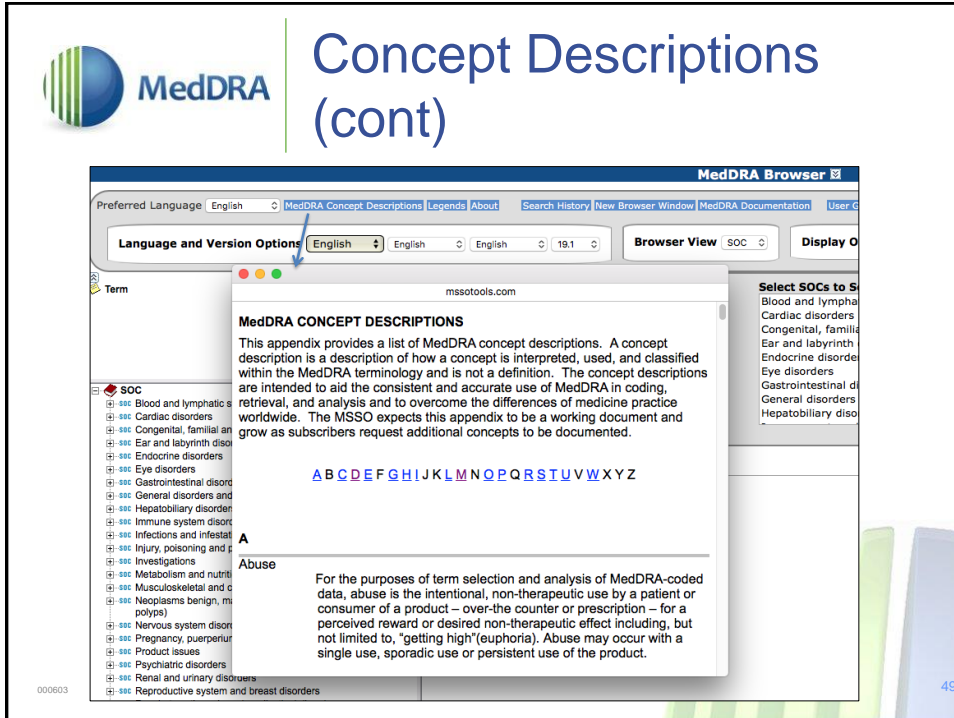
MedDRA

Concept Descriptions

- Descriptions of how a concept is interpreted, used, and classified in MedDRA
- Not a definition but important to consider when selecting MedDRA LLTs
- Intended to aid accurate and consistent use of MedDRA in coding and retrieval
- Overcome differences in medical practice worldwide
 - Descriptions aim to be broadly consistent with definitions across different regulatory regions
- Accessible in MSSO's browsers

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MedDRA Concept Descriptions (cont)

MedDRA Browser

Preferred Language: English | MedDRA Concept Descriptions | Legends | About | Search History | New Browser Window | MedDRA Documentation | User C

Language and Version Options: English | English | English | 19.1 | Browser View: SOC | Display O

Term: mssotools.com

MedDRA CONCEPT DESCRIPTIONS

This appendix provides a list of MedDRA concept descriptions. A concept description is a description of how a concept is interpreted, used, and classified within the MedDRA terminology and is not a definition. The concept descriptions are intended to aid the consistent and accurate use of MedDRA in coding, retrieval, and analysis and to overcome the differences of medicine practice worldwide. The MSSO expects this appendix to be a working document and grow as subscribers request additional concepts to be documented.

AB C D E F G H I J K L M N O P Q R S T U V W X Y Z

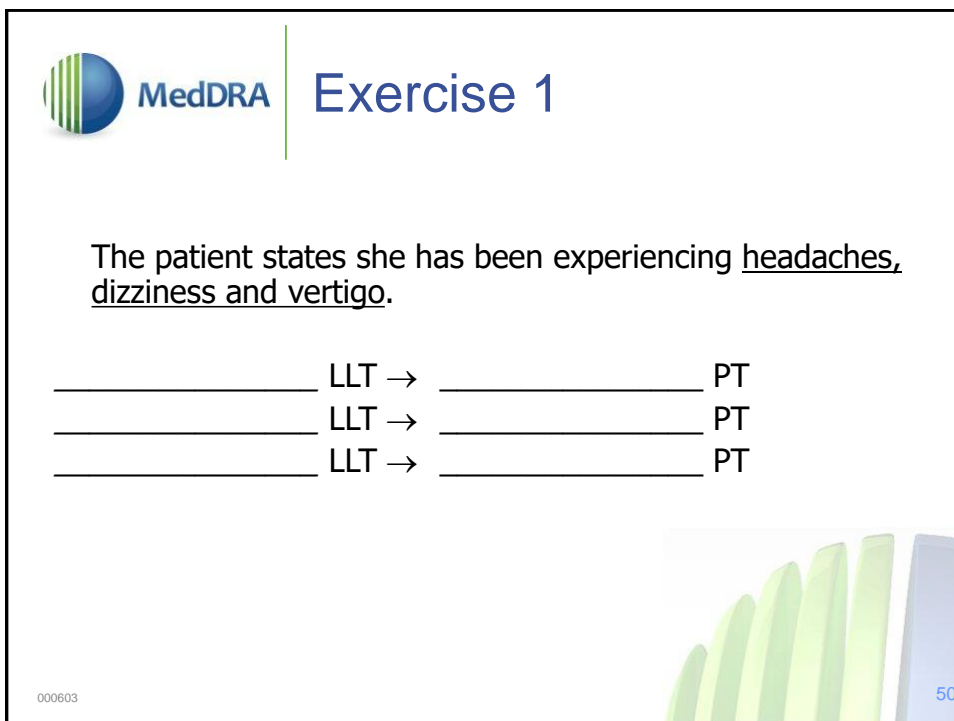
A

Abuse

For the purposes of term selection and analysis of MedDRA-coded data, abuse is the intentional, non-therapeutic use by a patient or consumer of a product – over-the counter or prescription – for a perceived reward or desired non-therapeutic effect including, but not limited to, "getting high"(euphoria). Abuse may occur with a single use, sporadic use or persistent use of the product.

Select SOCs to S: Blood and lymphatic, Cardiac disorders, Congenital, familial an, Ear and labyrinth, Endocrine disorder, Eye disorders, Gastrointestinal d, General disorders, Hepatobiliary diso

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MedDRA Exercise 1

The patient states she has been experiencing headaches, dizziness and vertigo.

_____ LLT → _____ PT

_____ LLT → _____ PT

_____ LLT → _____ PT

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MedDRA

Exercise 2

Lab results indicate an increase in erythrocytes.

_____ LLT → _____ PT

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MedDRA

Exercise 3

Drug was contaminated with Staphylococcus.

_____ LLT → _____ PT

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MedDRA

Exercise 4

A six year old boy was admitted for toxicity after accidentally ingesting the remaining antihypertensive tablets in the bottle.

_____ LLT → _____ PT

_____ LLT → _____ PT

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MedDRA

Exercise 5

The patient's urinary catheter was blocked.

_____ LLT → _____ PT

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Coding with MedDRA

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What are Coding Conventions?

- Written guidelines for coding with MedDRA in your organization
- Support accuracy and consistency
- Common topics
 - Misspellings, abbreviations and acronyms
 - Combination terms and "due to" concepts
 - "Always query" terms, e.g., "Chest pain"

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MedDRA

Why Do We Need Coding Conventions?

- Differences in medical aptitude of coders
- Consistency concerns (many more “choices” to manually code terms in MedDRA compared to older terminologies)
- Even with an autoencoder, may still need manual coding

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MedDRA

Autoencoder Pitfalls

- Inappropriate terms may be selected by autoencoder
- Review all autoencoding carefully
 - “Allergic to CAT scan” autoencoded as:
LLT *Allergic to cats*
 - “Myocardial infarction in the fall of 2000” autoencoded as:
LLT *Myocardial infarction*
LLT *Fall*

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MedDRA

Can I Make Coding Conventions Specific to My Company/Product?

- MedDRA may reduce the need to do this because:
 - Increased size/granularity results in more accurate representation of data
 - Secondary SOC allocations allow for different “views” of the data
- This type of approach should be done cautiously

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MedDRA

Synonym Lists

- Recurring verbatims – one-time assignment to an LLT
- Promotes consistency
- Increases likelihood of autoencoding “hit”
- Maintenance required

Verbatim	LLT	Comment
Throbbing above temple Aching all over head Pulsing pain in head	Headache	
Muscular pain in legs	Myalgia of lower extremities	LLT <i>Myalgia of lower extremities</i> is a better choice than LLT <i>Muscular pain</i> since it captures both the event and body site

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MedDRA

Quality Assurance (QA) Reports

- Allows reviewers to check for consistency (both auto-encoded and human-coded terms)
- Check for adherence to/deviation from coding conventions
- Check for emerging drifts/biases
- Multiple data views (verbatimim to coded terms; coded term to verbatimim; by SOC, etc.)

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MedDRA

QA Sample Report

SOC	HLT	PT	Verbatim	Count
Respiratory, thoracic and mediastinal disorders				
	Bronchospasm and obstruction			
		Wheezing		
			WHEEZING	16
			Wheeze	5
			INCREASED WHEEZING	1
			Breathing suppressed wheezing	1
			HYPERREACTIVITY AND WHEEZING	1
			wheeze in chest	1
	Laryngeal and adjacent sites disorders NEC (excl infections and neopla			
		Vocal cord disorder		
			SPASMODIC DYSTONIA OF THE VOCAL CORDS	1
	Newborn respiratory disorders NEC			
		Transient tachypnoea of the newborn		
			Transient hazy vision	1
			Transient tachypnea, neonatal	1
			Tachypnea of the newborn, transient	1

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MedDRA

Unqualified Test Name Term List

- MSSO developed and maintains list of unqualified test name terms
 - These terms (e.g., PT *Blood glucose*) should never be reported as AEs
 - Intended for use in E2B test name field only
- List can be used to check data quality
 - Identifies inappropriate terms in data fields other than test name data element
 - Intended as recommendation only

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Unqualified Test Name Term List (cont)

- Link on Support Documentation page on MedDRA website
- Spreadsheet of LLT/PT names and codes from SOC *Investigations*
 - >4,000 terms in v24.0
- Explanatory document
 - Purpose, uses, development of list
- Also available in Japanese on JMO website


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MedDRA Term Selection: Points to Consider Document

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MedDRA Term Selection: Points to Consider (MTS:PTC)

**MedDRA® TERM SELECTION:
POINTS TO CONSIDER**
ICH-Endorsed Guide for MedDRA Users

Release 4.21

March 2021

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- Provides term selection advice for industry and regulatory purposes
- Objective is to promote accurate and consistent term selection to facilitate a common understanding of shared data
- Recommended to be used as basis for individual organization's own coding conventions

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MedDRA

MedDRA Term Selection: PTC (cont)

- Developed by a working group of the ICH Management Committee
- Updated annually in step with the March release of MedDRA
- Complete versions available in English, Japanese, Chinese, Korean, and Spanish
- Condensed versions available for other MedDRA languages
- Available on MedDRA and JMO websites

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MedDRA

MTS:PTC Points of Note

- In some cases, with more than one option for selecting terms, a “preferred option” is identified but this does not limit MedDRA users to applying that option. Organizations should be consistent in their choice of option.
- Section 4.1 – Versioning (Appendix)
 - 4.1.1 Versioning methodologies
 - 4.1.2 Timing of version implementation

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MedDRA

PtC Documents

PtC Category	PtC Document	Purpose	Languages	Release Cycle
Term Selection	MedDRA Term Selection: Points to Consider	Promote accurate and consistent coding with MedDRA	English, Japanese, Chinese, Korean, and Spanish	Updated annually with the March release of MedDRA
	MedDRA Term Selection: Points to Consider Condensed Version	Shorter version focusing on general coding principles to promote accurate and consistent use of MedDRA worldwide	All MedDRA languages (except English, Japanese, and other languages with an available translation of the full MTS:PTC document)	Update as needed
Data Retrieval and Presentation	MedDRA Data Retrieval and Presentation: Points to Consider	Demonstrate how data retrieval options impact the accuracy and consistency of data output	English, Japanese, Chinese, Korean, and Spanish	Updated annually with the March release of MedDRA
	MedDRA Data Retrieval and Presentation: Points to Consider Condensed Version	Shorter version focusing on general retrieval and analysis principles to promote accurate and consistent use of MedDRA worldwide	All MedDRA languages (except English, Japanese, and other languages with an available translation of the full DRP:PTC document)	Update as needed

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PtC Documents (cont)

PtC Category	PtC Document	Purpose	Languages	Release Cycle
General	MedDRA Points to Consider Companion Document	More detailed information, examples, and guidance on specific topics of regulatory importance. Intended as a "living" document with updates based on users' needs. First edition covered data quality and medication errors. Updated in July 2020 (Release 1.1). Release 2.0 in October 2020 included new section on product quality issues. Another new section on manufacturing product quality issues is being drafted.	English and Japanese	Updated as needed

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General Term Selection Principles

- Quality of Source Data
- Quality Assurance
- Do Not Alter MedDRA
- Always Select a Lowest Level Term
- Select Only Current Lowest Level Terms
- When to Request a Term
- Use of Medical Judgment in Term Selection
- Select Terms for All Reported Information, Do Not Add Information

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Quality of Source Data Quality Assurance

- Quality of original information impacts quality of output
- Obtain clarification of data
- Can be optimized by careful design of data collection forms and proper training of staff
- Organizations' coding guidelines should be consistent with MTS:PTC
- Review of term selection by qualified individuals
- Human oversight of automated coding results

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Do Not Alter MedDRA

- MedDRA is a standardized terminology with a pre-defined term hierarchy
- Users must not make *ad hoc* structural alterations, including changing the primary SOC allocation



If terms are incorrectly placed, submit a change request to the MSSO

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Always Select a Lowest Level Term Select Only Current LLTs

- Lowest Level Term that most accurately reflects the reported verbatim information should be selected
- Degree of specificity may be challenging
 - Example: "*Abscess on face*" → select "*Facial abscess*," not simply "*Abscess*"
- Select current LLTs only
 - Non-current terms for legacy conversion/historical purposes

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When to Request a Term Use of Medical Judgment

- Avoid company-specific “work-arounds” for MedDRA deficiencies. If concept not adequately represented in MedDRA, submit Change Request to MSSO.
- If no exact match in MedDRA, use medical judgment to match to an existing term that adequately represents the concept

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Select Terms for All Reported Information

- Select terms for every AR/AE reported, regardless of causal association
- Select terms for device-related events, product quality issues, medication errors, medical and social history, investigations and indications as appropriate

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MedDRA

Do Not Add Information

- Do not make diagnosis if only signs/symptoms reported

Reported	LLT Selected	Comment
Abdominal pain, increased serum amylase, and increased serum lipase	Abdominal pain	It is inappropriate to assign an LLT for diagnosis of "pancreatitis"
	Serum amylase increased	
	Lipase increased	

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Selecting More than One Term Check the Hierarchy

- Can select more than one LLT to represent reported information. Document procedures.
 - Selecting one term may lead to loss of specificity
 - Selecting more than one term may lead to redundant counts
- Check the hierarchy above a selected LLT (PT, HLT, HLGT, SOC) to ensure placement accurately reflects meaning of reported term

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MedDRA

Term Selection Points

- Diagnoses and Provisional Diagnoses with or without Signs and Symptoms
- Death and Other Patient Outcomes
- Suicide and Self-Harm
- Conflicting/Ambiguous/Vague Information
- Combination Terms
- Age vs. Event Specificity
- Body Site vs. Event Specificity
- Location-Specific vs. Microorganism-Specific Infection
- Modification of Pre-existing Conditions
- Exposures During Pregnancy and Breast Feeding
- Congenital Terms
- Neoplasms
- Medical and Surgical Procedures
- Investigations

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Term Selection Points (cont)

- Medication Errors, Accidental Exposures and Occupational Exposures
- Misuse, Abuse and Addiction
- Transmission of Infectious Agent via Product
- Overdose, Toxicity and Poisoning
- Device-related Terms
- Drug Interactions
- No Adverse Effect and "Normal" Terms
- Unexpected Therapeutic Effect
- Modification of Effect
- Social Circumstances
- Medical and Social History
- Indication for Product Use
- Off Label Use
- Product Quality Issues

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Important Coding Errors

- Missed Concepts
 - All medical concepts described after the product is taken should be coded
 - Example: “*The patient took drug X and developed alopecia, increased LFTs and pancreatitis*”. Manufacturer only codes alopecia and increased LFTs (missed concept of pancreatitis)
 - Example: “*The patient took drug X and developed interstitial nephritis which later deteriorated into renal failure*”. Manufacturer only codes interstitial nephritis (missed renal failure concept)

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Important Coding Errors (cont)

- “Soft Coding”
 - Selecting a term which is both less specific and less severe than another MedDRA term is “soft coding”
 - Example: “*Liver failure*” coded as hepatotoxicity or increased LFTs
 - Example: “*Aplastic anemia*” coded as unspecified anemia
 - Example: “*Rash subsequently diagnosed as Stevens Johnson syndrome*” coded as rash

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Diagnoses and Provisional Diagnoses

SINGLE DIAGNOSIS	
DEFINITIVE DIAGNOSIS	PROVISIONAL DIAGNOSIS
Single diagnosis without signs and symptoms •Diagnosis (only possible option)	Single provisional diagnosis without signs and symptoms •Provisional diagnosis (only possible option)
Example: "Myocardial infarction" → select "Myocardial infarction"	Example: "Possible myocardial infarction" → select "Myocardial infarction" (select term as if definitive diagnosis)

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Similar principles apply for multiple diagnoses

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Diagnoses and Provisional Diagnoses (cont)

SINGLE DIAGNOSIS	
DEFINITIVE DIAGNOSIS	PROVISIONAL DIAGNOSIS
Single diagnosis with signs/symptoms •Preferred: Diagnosis only	Single provisional diagnosis with signs/symptoms •Preferred: Provisional diagnosis and signs/symptoms
Example: "Anaphylactic reaction with rash, dyspnoea, hypotension, and laryngospasm" → select "Anaphylactic reaction"	Example: "Possible myocardial infarction with chest pain, dyspnoea, diaphoresis" → select "Myocardial infarction", "Chest pain", "Dyspnoea", and "Diaphoresis"

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Similar principles apply for multiple diagnoses

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Diagnoses and Provisional Diagnoses (cont)

SINGLE DIAGNOSIS	
DEFINITIVE DIAGNOSIS	PROVISIONAL DIAGNOSIS
Single diagnosis with signs/symptoms •Alternate: Diagnosis and signs/symptoms	Single provisional diagnosis with signs/symptoms •Alternate: Signs/symptoms only (as provisional diagnosis may change)
Example: "Anaphylactic reaction with rash, dyspnoea, hypotension, and laryngospasm" → select "Anaphylactic reaction", "Rash", "Dyspnoea", "Hypotension", and "Laryngospasm"	Example: "Possible myocardial infarction with chest pain, dyspnoea, diaphoresis" → select "Chest pain", "Dyspnoea", and "Diaphoresis"

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Similar principles apply for multiple diagnoses

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Diagnoses and Provisional Diagnoses (cont)

- Always include signs/symptoms not associated with diagnosis

Reported	LLT Selected
Myocardial infarction, chest pain, dyspnoea, diaphoresis, ECG changes and jaundice	Myocardial infarction Jaundice (note that jaundice is not typically associated with myocardial infarction)

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What Terms to Select?

- Sepsis leading to shock from possible spontaneous bacterial peritonitis or bowel perforation

Sepsis

Shock

Septic shock

Spontaneous bacterial peritonitis

Bowel perforation

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MedDRA

Conflicting/Ambiguous Information

- First, try to obtain more specific information

Reported	LLT Selected	Comment
Hyperkalaemia with a serum potassium of 1.6 mEq/L	Serum potassium abnormal	LLT <i>Serum potassium abnormal</i> covers both of the reported concepts (note: serum potassium of 1.6 mEq/L is a low result, not high)
GU pain	Pain	"GU" could be either "genito-urinary" or "gastric ulcer". If additional information is not available, then select a term to reflect the information that is known, i.e., LLT <i>Pain</i>

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Vague Information

- First, try to obtain more specific information

Reported	LLT Selected	Comment
Turned green	Unevaluable event	"Turned green" reported alone is vague; this could refer to a patient condition or even to a product (e.g., pills)
Patient had a medical problem of unclear type	Ill-defined disorder	Since it is known that there is some form of a medical disorder, LLT <i>Ill-defined disorder</i> can be selected

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MedDRA

What Terms to Select?

- Clinical complication of IUD

IUD complication (PT Complication associated with device)

Intra-uterine death (PT Foetal death)

Unevaluable event

- Hypoglycemia (blood glucose = 200 mg/dL)

Blood glucose abnormal

Blood glucose increased

Hypoglycemia

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Combination Terms

- One condition is more specific than the other

Reported	LLT Selected
Arrhythmia due to atrial fibrillation	Atrial fibrillation
Hepatic function disorder (acute hepatitis)	Hepatitis acute

- A MedDRA combination term is available

Reported	LLT Selected
Retinopathy due to diabetes	Diabetic retinopathy
Rash with itching	Itchy rash
Breast cancer (HER2 positive)	HER2 positive breast cancer

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MedDRA

Combination Terms (cont)

- If splitting provides more clinical information, select more than one term
- In all cases of combination terms, apply medical judgment

Reported	LLT Selected
Diarrhoea and vomiting	Diarrhoea Vomiting
Wrist fracture due to fall	Wrist fracture Fall
BRAF positive malignant melanoma	BRAF gene mutation Malignant melanoma

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MedDRA

What Terms to Select?

- Retinal disease from HIV with near total blindness (R and L)

Retinal damage

Retinal disorder

HIV disease

Blindness

HIV retinopathy

Blindness, both eyes

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MedDRA

Investigations

- Medical condition vs. investigation result

Reported	LLT Selected	Comment
Hypoglycaemia	Hypoglycaemia	LLT <i>Hypoglycaemia</i> links to SOC <i>Metabolism and nutrition disorders</i>
Decreased glucose	Glucose decreased	LLT <i>Glucose decreased</i> links to SOC <i>Investigations</i>

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Investigations (cont)

- Unambiguous investigation result

Reported	LLT Selected	Comment
Glucose 40 mg/dL	Glucose low	Glucose is clearly below the reference range

- Ambiguous investigation result

Reported	LLT Selected	Comment
His glucose was 40	Glucose abnormal	No units have been reported. Select LLT <i>Glucose abnormal</i> if clarification cannot be obtained.

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Investigations (cont)

- Investigation results consistent with diagnosis

Reported	LLT Selected	Comment
Elevated potassium, K 7.0 mmol/L, and hyperkalaemia	Hyperkalaemia	It is not necessary to select LLT <i>Potassium increased</i>

- Grouped investigation result terms

Reported	LLT Selected	Comment
Increased alkaline phosphatase, increased SGPT, increased SGOT and elevated LDH	Alkaline phosphatase increased SGPT increased SGOT increased LDH increased	Select four individual terms. A single term such as LLT <i>Liver function tests abnormal</i> should not be selected.

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What Terms to Select?

- Testing showed increased serum creatinine and BUN, with increased BUN/creatinine ratio

Increased serum creatinine

BUN increased

Blood urea nitrogen/creatinine ratio increased

Renal function tests NOS abnormal

- Patient had features of aldosterone excess

Aldosterone increased

Aldosteronism

Blood aldosterone abnormal

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Medication Errors

- See MedDRA browsers (both WBB and MDB) for Concept Descriptions related to medication errors
- See Section 3 of MedDRA Points to Consider Companion Document for detailed examples, guidance and "Questions and Answers" about medication errors
- "Top-down" navigation in HLGT *Medication errors and other product use errors and issues* is best approach for term selection

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Medication Errors (cont)

- Medication error with clinical consequences

Reported	LLT Selected	Comment
Patient was administered wrong drug and experienced hypotension	Wrong drug administered Hypotension	
Insulin was given using the wrong syringe resulting in the administration of an overdose. The patient developed hypoglycaemia.	Drug administered in wrong device Accidental overdose Hypoglycaemia	If an overdose is reported in the context of a medication error, the more specific term <i>LLT Accidental overdose</i> can be selected

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Medication Errors (cont)

- Medication error without clinical consequences

Reported	LLT Selected	Preferred Option
Medication was given intravenously instead of intramuscularly without any adverse effect	Intramuscular formulation administered by other route	✓
	Intramuscular formulation administered by other route No adverse effect	

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Medication Errors (cont)

- Important to record potential occurrence of medication error
- Unlikely to be reported as an adverse event but may need to be recorded in periodic safety reports

Reported	LLT Selected	Comment
Pharmacist notices that the names of two drugs look similar and is concerned that this may result in a medication error	Drug name look-alike Circumstance or information capable of leading to medication error	Note: this example is a potential medication error. LLT <i>Drug name look-alike</i> provides additional information about the nature of the potential medication error, and LLT <i>Circumstance or information capable of leading to medication error</i> indicates that there is a potential medication error.

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Overdose, Toxicity and Poisoning

If overdose, poisoning or toxicity is explicitly reported, select the appropriate term

- Overdose with clinical consequences

Reported	LLT Selected
Stomach upset from study drug overdose	Stomach upset Overdose

- Overdose without clinical consequences

Reported	LLT Selected	Preferred Option
Patient received an overdose of medicine without any adverse consequences	Overdose	✓
	Overdose No adverse effect	

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What Term(s) to Select?

- The patient's renal function was measured every six months instead of on the monthly schedule recommended in the label for the drug

Medication monitoring error

Renal function test abnormal

Drug monitoring procedure incorrectly performed

- Unintentionally took more than maximum recommended dose due to dispensing error

Accidental overdose

Incorrect dose administered

Drug dispensing error

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Death and Other Patient Outcomes

- Death, disability and hospitalization are considered outcomes and not usually coded as adverse events (AE) or adverse reactions (AR)
- If one of these outcomes is the only reported information, select an appropriate LLT for coding
- If a death term in MedDRA adds important clinical information, code it along with reported ARs/AEs

Reported	LLT Selected
Patient experienced a rash and had sudden cardiac death	Rash Sudden cardiac death

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Drug Interactions

- Term includes reactions between drugs (including biologic products) and other drugs, foods, devices and alcohol
- If reporter states an interaction, select terms for interaction term and reported medical event(s)

Reported	LLT Selected
Patient drank cranberry juice which interacted with anticoagulant drug causing an INR increase	Food interaction INR increased

- If reporter does not specifically state an interaction, select terms for reported medical event(s) only

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Suicide and Self-Harm

- Coding of suicide attempts, completed suicides and self-harm should be accurate and consistent
- Do not assume that an overdose, even if intentional, is a suicide attempt
- If a self-harm report does not mention suicide or suicide attempt, only code self-injury term
- If a suicide attempt is fatal, select a term that reflects the outcome instead of the attempt

Reported	LLT Selected	Comment
Suicide attempt resulted in death	Completed suicide	Report death as an outcome

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What Term to Select?

- Death from cerebral hemorrhage

Sudden death

Death

Cerebral hemorrhage

Brain death

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What Term to Select?

- Patient was found dead

Death from natural causes

Death

Died in sleep

Found dead

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What Term(s) to Select?

- After taking an antihistamine along with her prescription proton pump inhibitor, a 53-year-old woman developed vertigo

Drug interaction NOS

Vertigo subjective

Vertigo

Drug interaction

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What Term(s) to Select?

- Deliberately took an overdose

Intentional overdose

Suicide attempt

Deliberate overdose

Overdose NOS

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MedDRA Misuse, Abuse and Addiction

Concept	Intentional?	By Whom?	Therapeutic Use?	Additional Sections in this Document
Misuse	Yes	Patient/consumer	Yes*	3.16.1
Abuse	Yes	Patient/consumer	No	3.16.2
Addiction	Yes	Patient/consumer	No	3.16.3
Medication error	No	Patient/consumer or healthcare provider	Yes	3.15
Off label use	Yes	Healthcare provider	Yes	3.27

* Definitions of misuse may not always include the concept of therapeutic use; misuse may be similar to the concept of abuse in some regions.

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MedDRA Coding Exercises

- Narratives and short verbatims
- Assess the reported terms
 - Identify what concepts are reported (diagnosis, death, investigations, etc.)
- Refer to the appropriate sections of the MTS:PTC for guidance on term selection
 - For example, Section 3.2 for death terms
- Use MTS:PTC preferred options (forget your organization's conventions)
- Use browser to search for and select LLTs

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Specific Tips for Narrative Exercises

- Overall, coding principles are the same as for short verbatim exercises
- Code all of the following:
 - Events (including procedures and investigations as needed)
 - Indications
 - Medical history
 - Social history

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Sample Narrative

A 75-year-old male receiving Drug X for rheumatoid arthritis developed an area of darkened skin on his chest. The patient's medical history is significant for peripheral vascular disease and cigarette smoking. The skin lesion was excised; it was revealed to be a seborrhoeic wart.

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Course Summary

- In this course, we covered:
 - An overview of MedDRA's background, scope, and structure (including primary SOC allocation rules)
 - Maintenance of MedDRA
 - Coding conventions, synonym lists, and coding QA
 - Introduction to the MedDRA Term Selection: Points to Consider document
 - Coding exercises

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MSSO Contacts

- Website
 - www.meddra.org
- Email
 - mssohelp@meddra.org
- Frequently Asked Questions
 - www.meddra.org/faq
- MedDRA Browsers
 - <https://www.meddra.org/meddra-desktop-browsers> (Desktop Browser)
 - <https://tools.meddra.org/wbb/> (Web-Based Browser)
 - <https://mmb.meddra.org> (Mobile Browser)

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MSSO Contacts (cont)


- Change Request Submission
 - <https://www.meddra.org/how-to-use/change-requests>
- Training Schedule
 - <https://www.meddra.org/training/schedule>
- MedDRA Support Documentation
 - <https://www.meddra.org/how-to-use/support-documentation>

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Thank You!



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Questions ?

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