Getting Started with MedDRA

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).
Disclaimer and Copyright Notice

- This presentation is protected by copyright and may, with the exception of the MedDRA and ICH logos, be used, reproduced, incorporated into other works, adapted, modified, translated or distributed under a public license provided that ICH's copyright in the presentation is acknowledged at all times. In case of any adaption, modification or translation of the presentation, reasonable steps must be taken to clearly label, demarcate or otherwise identify that changes were made to or based on the original presentation. Any impression that the adaption, modification or translation of the original presentation is endorsed or sponsored by the ICH must be avoided.
- The presentation is provided "as is" without warranty of any kind. In no event shall the ICH or the authors of the original presentation be liable for any claim, damages or other liability arising from the use of the presentation.
- The above-mentioned permissions do not apply to content supplied by third parties. Therefore, for documents where the copyright vests in a third party, permission for reproduction must be obtained from this copyright holder.

Course Overview

- Describe MedDRA governance
- Discuss MedDRA's background, scope, structure, and characteristics
- Discuss MedDRA implementation topics
- Describe need for procedures documentation and conventions
- Describe Standardised MedDRA Queries (SMQs)
- Highlight MedDRA tools and training
- Conclude with a question and answer session
MedDRA Governance

Governance Structure for MedDRA

- ICH MedDRA Management Committee appointed by the ICH Assembly to provide oversight of MedDRA related activities and the Maintenance and Support Services Organization (MSSO)
Management Committee/MSSO Relationship

- ICH owns MedDRA
- ICH MedDRA Management Committee
  - Contracts with MSSO to maintain it
  - Has oversight of all operations of the MSSO
    - Meets regularly with MSSO
    - Sets subscription rates
    - Approves developmental plans and services
  - Membership includes ICH regulatory authorities and industry associations

MedDRA Background, Scope, Structure, and Characteristics
What is MedDRA?

Med = Medical
D = Dictionary for
R = Regulatory
A = Activities

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
Where MedDRA is Used

Preclinical Testing ➤ Clinical Phase I ➤ Clinical Phase II ➤ Clinical Phase III ➤ Marketed Product Phase IV

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

Electronic Transmission of Data

INTERNATIONAL CONFERENCE ON HARMONISATION OF TECHNICAL REQUIREMENTS FOR REGISTRATION OF PHARMACEUTICALS FOR HUMAN USE

ICH E2B Expert Working Group

Implementation Guide for
Electronic Transmission of Individual Case Safety Reports (ICSRs)

E2B(R3) Data Elements and Message Specification

Version 5.01, 12 April 2013
• As of January 2019
  – 5,700 Subscribing organizations (MSSO+JMO)
  – 122 Countries
• Graph shows types of subscribing organizations

MedDRA Users by Region

<table>
<thead>
<tr>
<th>Country</th>
<th>Count</th>
</tr>
</thead>
<tbody>
<tr>
<td>United States</td>
<td>1575</td>
</tr>
<tr>
<td>Japan</td>
<td>785</td>
</tr>
<tr>
<td>UK</td>
<td>336</td>
</tr>
<tr>
<td>Germany</td>
<td>325</td>
</tr>
<tr>
<td>China</td>
<td>299</td>
</tr>
<tr>
<td>France</td>
<td>245</td>
</tr>
<tr>
<td>Italy</td>
<td>207</td>
</tr>
<tr>
<td>Spain</td>
<td>150</td>
</tr>
<tr>
<td>Canada</td>
<td>127</td>
</tr>
<tr>
<td>Republic of Korea</td>
<td>111</td>
</tr>
<tr>
<td>Sweden</td>
<td>96</td>
</tr>
<tr>
<td>Australia</td>
<td>95</td>
</tr>
<tr>
<td>Netherlands</td>
<td>92</td>
</tr>
<tr>
<td>India</td>
<td>90</td>
</tr>
<tr>
<td>Switzerland</td>
<td>84</td>
</tr>
<tr>
<td>Poland</td>
<td>71</td>
</tr>
<tr>
<td>Belgium</td>
<td>61</td>
</tr>
<tr>
<td>Chinese Taipei</td>
<td>60</td>
</tr>
<tr>
<td>Israel</td>
<td>57</td>
</tr>
<tr>
<td>Greece</td>
<td>56</td>
</tr>
<tr>
<td>Portugal</td>
<td>52</td>
</tr>
<tr>
<td>Denmark</td>
<td>50</td>
</tr>
<tr>
<td>Austria</td>
<td>43</td>
</tr>
<tr>
<td>Russian Federation</td>
<td>37</td>
</tr>
<tr>
<td>Czechia</td>
<td>36</td>
</tr>
</tbody>
</table>
MedDRA Data Sharing

- Subscription grants access to MedDRA for one year
- Subscriber cannot grant any sublicense, publish or otherwise distribute MedDRA to a third party
- Data may be freely exchanged between current MedDRA subscribers
  - Sponsor-sponsor, sponsor-CRO, vendor-user, etc.
  - Use Self-Service Application to check organization’s subscription status
- Sharing MedDRA with a non-subscribing organization is a violation of the MedDRA license

MedDRA Data Sharing (cont)

- For details, see the Statement on MedDRA Data Sharing
Scope of MedDRA

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

OUT

Not a drug dictionary
Frequency qualifiers
Numerical values for results
Severity descriptors
Not an equipment, device, diagnostic product dictionary

Patient demographic terms
Clinical trial study design terms

MedDRA Structure

System Organ Class (SOC) (27)
High Level Group Term (HLGT) (337)
High Level Term (HLT) (1,737)
Preferred Term (PT) (23,708)
Lowest Level Term (LLT) (80,262)
System Organ Classes

- 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
- Infections and infestations
- Injury, poisoning and procedural complications
- Investigations
- Metabolism and nutrition disorders
- Musculoskeletal and connective tissue disorders
- Neoplasms benign, malignant and unspecified (incl cysts and polyps)
- Nervous system disorders
- Pregnancy, puerperium and perinatal conditions
- Product issues
- Psychiatric disorders
- Renal and urinary disorders
- Reproductive system and breast disorders
- Respiratory, thoracic and mediastinal disorders
- Skin and subcutaneous tissue disorders
- Social circumstances
- Surgical and medical procedures
- Vascular disorders

Lowest Level Term

Synonyms, lexical variants, sub-elements

SOC = Cardiac disorders

HLGT = Cardiac arrhythmias

HLT = Rate and rhythm disorders NEC

PT = Arrhythmia

LLT = Arrhythmia

LLT (Non-current) Other specified cardiac dysrhythmias

Not all LLTs shown
Non-Current Terms

- 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

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
A Multi-Axial Terminology

- Multi-axial = the representation of a medical concept in multiple SOCs
  - 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
A Multi-Axial Terminology (cont)

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

**HLGT** = Respiratory tract infections

**HLT** = Viral upper respiratory tract infections

**PT** = Influenza

**SOC** = Infections and infestations (Primary SOC)

**HLGT** = Viral infectious disorders

**HLT** = Influenza viral infections

MSSO’s MedDRA Browsers

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

- **MedDRA Web-Based Browser (WBB)**
  - [https://tools.meddra.org/wbb/](https://tools.meddra.org/wbb/)

**Features**

- Both 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
MedDRA Desktop Browser (MDB) and Web-Based Browser (WBB) Update

• New functionality for users
  – 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

MedDRA Browser SOC View
MedDRA Implementation Topics

Implementing MedDRA

- No single method for all organizations
- Implementation approach depends on factors such as size of organization
**Downloading and Installing MedDRA**

- Step-by-step instructions in Appendix slides
  - Downloading the latest release of MedDRA
  - Installing the MedDRA Desktop Browser
  - Loading MedDRA into the Desktop Browser

**MedDRA Maintenance**

- MSSO processes change requests from its subscriber organizations
- Rigorous medical review by MSSO physicians
- Twice yearly official updates
  - 1 March X.0 release
  - 1 September X.1 release
Reasons to Update to a New MedDRA Version

• Take advantage of new terms and other improvements
• Use the same new version to summarize data from different sources that had used older versions
  – “Pooling” clinical trials for analysis
  – Post-marketing safety summaries, etc.
• Use of recent versions of MedDRA may be required or preferred by regulatory authorities

Reasons to Update to a New MedDRA Version (cont)

• Stay current with development partners and contract research organizations
• Harmonize use of MedDRA to optimize communication of data
Resources for Version Updates

- MedDRA “Best Practices” document
- Appendix 4.1 Versioning in the MedDRA Term Selection: Points to Consider document
- MSSO videocast – MedDRA Version Updates
- What’s New document
- MSSO “What’s New” webinars
- MedDRA Version Reports
- MedDRA Version Analysis Tool (MVAT)

Version Report

- List of various types of changes in MedDRA
MedDRA Version Analysis Tool

Procedures Documentation and Conventions
Procedures Documentation

- Documentation of procedures and processes is a best practice for all organizations
  - This includes the use of MedDRA
- Standard operating procedures and other relevant documents in an organization should address the use of MedDRA
  - Coding, dictionary management, MedDRA versioning, and analytical processes and tools (e.g., using MedDRA’s hierarchy and Standardised MedDRA Queries)

Need for Conventions

- Conventions are written guides and sets of principles for using MedDRA
- MedDRA may appear complicated but its structure is logical, and several guides are available to optimize its use
- Conventions help users achieve consistency in data entry (coding) and data retrieval
- Harmonizes exchange of MedDRA coded data worldwide
ICH M1 Points to Consider Working Group (PtC WG)

- Regulators and industry from EU, US, and Japan
- Health Canada, Canada
- MFDS, Republic of Korea
- ANVISA, Brazil
- NMPA, China
- MSSO
- JMO
- WHO (Observer)

November 2017, Geneva, Switzerland

PtC Documents

<table>
<thead>
<tr>
<th>PtC Category</th>
<th>PtC Document</th>
<th>Purpose</th>
<th>Languages</th>
<th>Release Cycle</th>
</tr>
</thead>
<tbody>
<tr>
<td>Term Selection</td>
<td>MedDRA Term Selection: Points to Consider</td>
<td>Promote accurate and consistent coding with MedDRA</td>
<td>English and Japanese</td>
<td>Updated with each MedDRA release</td>
</tr>
<tr>
<td></td>
<td>MedDRA Term Selection: Points to Consider Condensed Version</td>
<td>Shorter version focusing on general coding principles to promote accurate and consistent use of MedDRA worldwide</td>
<td>All MedDRA languages (except English and Japanese)</td>
<td>Update as needed</td>
</tr>
<tr>
<td>Data Retrieval and Presentation</td>
<td>MedDRA Data Retrieval and Presentation: Points to Consider</td>
<td>Demonstrate how data retrieval options impact the accuracy and consistency of data output</td>
<td>English and Japanese</td>
<td>Updated with each MedDRA release</td>
</tr>
<tr>
<td></td>
<td>MedDRA Data Retrieval and Presentation: Points to Consider Condensed Version</td>
<td>Shorter version focusing on general retrieval and analysis principles to promote accurate and consistent use of MedDRA worldwide</td>
<td>All MedDRA languages (except English and Japanese)</td>
<td>Update as needed</td>
</tr>
<tr>
<td>General</td>
<td>MedDRA Points to Consider Companion Document</td>
<td>More detailed information, examples, and guidance on specific topics of regulatory importance. Intended as a &quot;living&quot; document with frequent updates based on users' needs. First edition covers data quality and medication errors.</td>
<td>English and Japanese</td>
<td>Updated as needed</td>
</tr>
</tbody>
</table>
**MedDRA Term Selection: Points to Consider (MTS:PTC)**

- 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

---

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

**Release 4.17**
Based on MedDRA Version 22.0

1 March 2019

Disclaimer and Copyright Notice
This document is protected by copyright and may, with the exception of the MedDRA and ICH logos, be used, reproduced, incorporated into other works, adapted, modified, translated or distributed under a public license provided that ICH’s copyright in the document is acknowledged at all times. In case of any adoption, modification or translation of the document, reasonable steps must be taken to clearly label, demarcate or otherwise identify that changes were made to or based on the original document. Any impression that the adoption, modification or translation of the original document is endorsed or sponsored by the ICH must be avoided.
The document is provided "as is" without warranty of any kind. In no event shall the ICH or the authors of the original document be liable for any claim, damages or other liability arising from the use of the document.
The above-mentioned permissions do not apply to content supplied by third parties. Therefore, for documents where the copyright vests in a third party, permission for reproduction must be obtained from this copyright holder.
MedDRA® trademark is registered by IFPMA on behalf of ICH.

---

**MedDRA Term Selection: PTC (cont)**

- Developed by a working group of the ICH Management Committee
- Updated twice yearly with each MedDRA release
- Available on MedDRA and JMO websites
  - English and Japanese
  - Word ("clean" and "redlined"), PDF, HTML formats
  - "Redlined" document identifies changes made from previous to current release of document
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

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
**Coding: Translating into MedDRA**

<table>
<thead>
<tr>
<th>Reported Information</th>
<th>LLT</th>
</tr>
</thead>
<tbody>
<tr>
<td>Throbbing above temple</td>
<td>Headache</td>
</tr>
<tr>
<td>Aching all over head</td>
<td></td>
</tr>
<tr>
<td>Pulsing pain in head</td>
<td></td>
</tr>
<tr>
<td>Really bad headache</td>
<td></td>
</tr>
<tr>
<td>Headache</td>
<td></td>
</tr>
<tr>
<td>Infection in lungs</td>
<td>Lung infection</td>
</tr>
<tr>
<td>Patient took Drug A instead of Drug B and experienced hypertension</td>
<td>Wrong drug administered Hypertension</td>
</tr>
</tbody>
</table>

**MedDRA Data Retrieval and Presentation: Points to Consider (DRP:PTC)**

- Provides data retrieval and presentation options for industry or regulatory purposes
- Most effective when used in conjunction with MedDRA Term Selection: PTC document
- Recommended to be used as basis for individual organization’s own data retrieval conventions
High quality data output is dependent on maintaining quality of original information reported by using consistent and appropriate term selection (Refer to MedDRA Term Selection: Points to Consider document)

Method of conversion of data into MedDRA might impact retrieval and presentation - legacy data conversion using verbatims or coded terms

Legacy Data Conversion

Method 1 - Data converted from legacy terminology terms to MedDRA

<table>
<thead>
<tr>
<th>Reported</th>
<th>Legacy Term</th>
<th>MedDRA Term</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gastrointestinal ischaemia</td>
<td>Gastrointestinal disorder</td>
<td>Gastrointestinal disorder</td>
</tr>
</tbody>
</table>

- Results reflect specificity of legacy terminology
- Benefits of greater specificity of MedDRA not attained
Legacy Data Conversion (cont)

- WHO Uppsala Monitoring Centre and the MSSO maintain a WHO-ART to MedDRA bridge to support the conversion of WHO-ART coded data to MedDRA

<table>
<thead>
<tr>
<th>Reported Legacy Term</th>
<th>MedDRA Term</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gastrointestinal ischaemia</td>
<td>Gastrointestinal disorder</td>
</tr>
</tbody>
</table>

Specificity of reported term reflected by MedDRA term
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

Standardised MedDRA Queries (SMQs)
Standardised MedDRA Queries (SMQs)

- Collaboration between CIOMS (Council for International Organizations of Medical Sciences) and ICH (MSSO)
- Groupings of terms from one or more MedDRA SOCs related to medical condition or area of interest
- Terms relate to signs/symptoms, diagnoses, syndromes, physical findings, laboratory and other test data, etc.
- Intended to aid in case identification

How to “Run” SMQs (cont)

Clinical Trial Database
Safety Database

Case
LLT 1
LLT 2
LLT 3

Query
SMQ
PT
LLT
LLT
PT
LLT
LLT

"Hit"
SMQs in Production - Examples

- As of Version 22.0, a total of 104 level 1 SMQs in production
  - Agranulocytosis
  - Anaphylactic reaction
  - Cerebrovascular disorders
  - Convulsions
  - Depression and suicide/self-injury
  - Hepatic disorders
  - Hypersensitivity
  - Ischaemic heart disease
  - Lack of efficacy/effect
  - Medication errors
  - Osteonecrosis
  - Peripheral neuropathy
  - Pregnancy and neonatal topics
  - Pseudomembranous colitis
  - Rhabdomyolysis/myopathy
  - Severe cutaneous adverse reactions
  - Systemic lupus erythematosus

SMQ Applications

- Clinical trials
  - Where safety profile is not fully established, use multiple SMQs on routine basis as screening tool
  - Selected SMQs to evaluate previously identified issue (pre-clinical data or class effect)

- Post-marketing
  - Selected SMQs to retrieve cases for suspected or known safety issue
  - Signal detection (multiple SMQs employed)
  - Single case alerts
  - Periodic reporting (aggregate cases for safety and other issues, e.g., lack of efficacy)
MedDRA Browser SMQ View

MedDRA Tools and Training
IT Considerations

- Software tools support the use of MedDRA
  - Several are free with MedDRA subscription
    - Two browsers (Desktop and Web-Based)
    - MedDRA Version Analysis Tool (MVAT)
  - Software tools need driven by the volume of data
    - With small amounts, users can use simple software tools (e.g., free MSSO browsers, spreadsheets)
    - Larger implementations may need commercial data management products
    - List of third-party software tools on MedDRA website

MedDRA Training Opportunities
– Available for Users

- Free Face-to-Face (F2F) training
  - Coding with MedDRA
  - Safety Data Analysis and Standardised MedDRA Queries
  - Getting Started with MedDRA

- Free webinars
  - Getting Started with MedDRA
  - MedDRA Overview
  - MedDRA Coding Basics
  - Advanced MedDRA Coding
  - Data Analysis and Query Building with MedDRA
  - Standardised MedDRA Queries
  - What’s New with MedDRA (with each MedDRA release)
MedDRA Training Opportunities
– Available to All

• Free resources on MedDRA website
  – Slides for all F2F courses and webinars
  – Short videocasts on MedDRA-related topics
    • Available in several languages
    • Can be downloaded or viewed directly on website
    • Help trainees prepare for F2F courses

• Webinars and videocasts available on new MedDRA MSSO YouTube Channel

More Resources for MedDRA Users

• MedDRA website
  – Help Desk
  – Subscriptions
  – News and Events
  – MedDRA Best Practices document
  – Points to Consider documents
  – Terminology downloads
  – Training
  – Tools
  – MedDRA publications
  – User group meetings
  – Expert meetings
Summary

In this course, we:
• Described MedDRA governance
• Discussed MedDRA’s background, scope, structure, and characteristics
• Discussed MedDRA implementation topics
• Described need for procedures documentation and conventions
• Described Standardised MedDRA Queries
• Highlighted MedDRA tools and training

MSSO Contacts

• Website
  – www.meddra.org
• Email
  – mssohelp@meddra.org
• Frequently Asked Questions
  – www.meddra.org/faq
Question and Answer Session

Appendix:
Downloading and Installing MedDRA
First Steps

- You just received your subscription letter from the MSSO, what’s next?
  - Download the latest release
- Identify the components and distribute accordingly
  - Install the MedDRA Desktop Browser
  - Load MedDRA into the Desktop Browser

Download the Latest Release

Click on Downloads
Download the Latest Release (cont)

- Enter Subscriber ID and password

Download the Latest Release (cont)

- Select the version and language(s)
  - Password protected zip file
- Save the zip file to your computer
Extract the Release Files

• Zip file structure and contents
  – Select a file to extract
  – Enter Unzip password from Welcome letter

What’s in the Zip File?

Full Release of MedDRA Data and relationships files
Changes in this release
Description of contents of zip file
Narrative discussion of changes in this release
SMQ Spreadsheet
SMQ Introductory Guide
Spreadsheet with lists of new and modified terms
Details of implemented changes
File Format Descriptions
MedDRA Introductory Guide
What to do with the Contents of the Zip File?

<table>
<thead>
<tr>
<th>Item</th>
<th>Who should receive it?</th>
</tr>
</thead>
<tbody>
<tr>
<td>MedAscii Directory</td>
<td>IT staff to load MedDRA</td>
</tr>
<tr>
<td>SEQAscii Directory</td>
<td>IT staff to review changes</td>
</tr>
<tr>
<td>Readme</td>
<td>All users</td>
</tr>
<tr>
<td>SMQ Spreadsheet</td>
<td>Those using SMQs</td>
</tr>
<tr>
<td>SMQ Introductory Guide</td>
<td>Those using SMQs</td>
</tr>
<tr>
<td>Version Report</td>
<td>Those interested in version changes</td>
</tr>
<tr>
<td>What’s New</td>
<td>All users</td>
</tr>
<tr>
<td>Detail Report</td>
<td>Those interested in version changes</td>
</tr>
<tr>
<td>Distribution File Format</td>
<td>IT staff to load MedDRA</td>
</tr>
<tr>
<td>MedDRA Introductory Guide</td>
<td>All users</td>
</tr>
</tbody>
</table>

Contents of MedASCII Directory

- MedDRA term and relationship files
- ASCII File format
Download and Install MedDRA Desktop Browser

1) Click on Downloads page from Home page
2) Log in (if not already)
3) Click on Desktop Browsers

Download and Install MedDRA Desktop Browser (cont)

Download installation zip file
Download installation instructions
Load MedDRA into Desktop Browser

• Install MedDRA Desktop Browser
  – Create a directory on your computer
  – Unzip the contents of the zip file downloaded to this directory
  – Double click the file MedDRABrowserWIN.exe to start the browser
    • For future use, right click the MedDRABrowserWIN.exe file and choose “Send to” and then “Desktop” and an icon will be placed on your desktop

Load MedDRA into Desktop Browser (cont)

• Use the browse button to identify the directory where the MedDRA ASCII files are stored

• Click Import
Load MedDRA into Desktop Browser (cont)

- The load process will display a progress bar and then the load process will complete.
- At the top of MedDRA Desktop Browser screen, select the version of MedDRA to display.

- MedDRA Desktop Browser is loaded and ready for use.