

# MedDRA<sup>®</sup> DATA RETRIEVAL AND PRESENTATION: POINTS TO CONSIDER

ICH-Endorsed Guide for MedDRA Users  
on Data Output

*Release 3.7*  
*Based on MedDRA Version 17.0*

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## Section 1 – INTRODUCTION

The **Medical Dictionary for Regulatory Activities** terminology (MedDRA) was designed for sharing regulatory information for human medical products. However, unless users achieve consistency in how they assign terms to verbatim reports of symptoms, signs, diseases, etc., and in methods for data retrieval and evaluation, use of MedDRA cannot have the desired harmonising effect in the exchange of coded data.

MedDRA is a large terminology with very specific (“granular”) terms called Lowest Level Terms (LLTs) that serve to accurately record the reporter’s words (verbatim term). LLTs are generally synonyms linked to their parent terms known as Preferred Terms (PTs). PTs are also relatively specific and large in number.

While a highly granular terminology such as MedDRA reduces the need for interpretation at data entry, it impacts the processes of data retrieval, sorting and presentation necessary for support of drug development, pharmacovigilance and risk management. The hierarchical structure of MedDRA facilitates data retrieval by providing grouping terms (High Level Terms [HLTs] and High Level Group Terms [HLGTs]) that aggregate the very specific terms used for coding into broader medical categories. MedDRA’s multiaxiality (assignment of a PT to more than one System Organ Class [SOC]) allows flexibility in data retrieval via primary and secondary paths. Whilst grouping terms and multiaxiality permit a reasonable first approach to data retrieval, the complexity of MedDRA requires guidance to optimise the results.

This *Data Retrieval and Presentation: Points to Consider* (DRP:PTC) document is an ICH-endorsed guide for MedDRA users. It is updated in step with new MedDRA versions and is a companion document to MedDRA. It was developed and is maintained by a working group charged by the ICH Steering Committee. The working group consists of regulatory and industry representatives of the European Union, Japan and the United States, as well as representatives from the Canadian regulatory authority, the MedDRA Maintenance and Support Services Organization (MSSO) and the Japanese Maintenance Organization (JMO). (See Appendix, Section 6.2 for list of members).

The principles described in this document are most effective when used in conjunction with the principles described in the *MedDRA Term Selection: Points to Consider* document for data entry (coding). This document provides data retrieval and presentation options for either industry or regulatory purposes. Although MedDRA includes some data retrieval tools, this document addresses data retrieval in a broader context.

Examples in this document are based on MedDRA Version 17.0; they are intended to facilitate reader understanding and are **not** intended to imply regulatory requirements.

Figures referenced in the text are found in the Appendix, Section 6.3.

## 1.1 – Objectives of this Document

The objective of the DRP:PTC document is to demonstrate how data retrieval options impact the accuracy and consistency of data output. For example, certain drugs or therapeutic areas may need a customised approach for data output. Options for data input described in the *MedDRA Term Selection: Points to Consider* document – or in organisation-specific coding guidelines – should also be taken into consideration.

Organisations are encouraged to document their data retrieval and output strategies, methods and quality assurance procedures in organisation-specific guidelines which should be consistent with this DRP:PTC document.

## 1.2 – Reasons to Use MedDRA

MedDRA is used to report adverse reaction/adverse event (AR/AE) terms in individual case reports – both on paper or electronically. Its structure allows for aggregation of those reported terms in medically meaningful groupings to facilitate analysis of safety data. MedDRA can also be used to list AR/AE data in reports (tables, line listings, etc), compute frequencies of similar AR/AEs, and capture and analyse related data such as product indications, investigations, and medical and social history.

## 1.3 – How to Use this Document

The principles described in this document apply to all data encoded with MedDRA with a focus on aggregated data. This document does not address the use of MedDRA for single case reporting, labeling, medical evaluation and statistical methodology.

This *Points to Consider* document aims to help all MedDRA users, since the MedDRA terminology itself contains no specific guidelines for its use. The document provides a framework to foster **consistent** use of MedDRA for data analysis and presentation for medically meaningful review and analysis of clinical data.

This document describes the features of MedDRA and highlights the impact of MedDRA's structure, rules and conventions on data output. Examples and options described in the document are not intended to communicate specific regulatory reporting requirements or address specific database issues. This

document cannot address every situation, therefore, medical judgment should always be applied.

The document is not a substitute for MedDRA training. It is essential for users to have knowledge of MedDRA's structure and content. For optimal use of MedDRA, one should refer to the MedDRA *Introductory Guide*, the *Introductory Guide for Standardised MedDRA Queries (SMQs)* (See Appendix, Section 6.1), and the *MedDRA Term Selection: Points to Consider* document).

## Section 2 – GENERAL PRINCIPLES

### 2.1 – Quality of Source Data

High quality data output occurs when the quality of the information originally reported is maintained with consistent and appropriate term selection. Organisations should pursue continuous oversight of data quality. Data quality issues are also addressed in the *MedDRA Term Selection: Points to Consider* document.

#### 2.1.1 Data conversion considerations

Give special consideration to the method used to convert data from other terminologies into MedDRA. The methods used can impact retrieval and presentation strategies.

- Method 1 – Data converted from legacy terminology terms to MedDRA
  - Results will reflect the specificity of the previous terminology
  - The benefits of the greater specificity of MedDRA are not attained

Example

Reported	Legacy Term	MedDRA Term
Bowel ischaemia	Gastrointestinal Disorder	Gastrointestinal disorder

- Method 2 – Data converted from the original reported terms (verbatim terms) to MedDRA terms

Example

Reported	Legacy Term	MedDRA Term
Bowel ischaemia	Gastrointestinal Disorder	Bowel ischaemia

Document the data conversion method used, including the date of the conversion.

### 2.1.2 Impact of data conversion method

Combining the two conversion methods described above can affect interpretation of data output.

Example

<b>Data Output with Combined Data Conversion Methods</b>
If data have been converted directly from legacy terminology terms to MedDRA terms (Method 1), and if newly acquired data are coded directly from reported terms to MedDRA, the resulting differences in specificity could make interpretation difficult.

When designing a search strategy, it may be useful to examine the **reported terms** for data converted using Method 1. If the search has been based on specific MedDRA terms, data previously coded to non-specific terms may be otherwise overlooked.

Example

<b>Impact of Method 1 Conversion on Search Strategy</b>
If searching with MedDRA PT <i>Bowel ischaemia</i> , cases of bowel ischaemia coded with the legacy term <i>Gastrointestinal disorder</i> would be missed. In this case, it would be important to know the date of the legacy data conversion.

To conduct a search requiring this level of detail, it might be necessary to review or recode from the reported terms. For legacy data, this information might be found in fields other than those for ARs/AEs.

## 2.2 – Documentation of Data Retrieval and Presentation Practices

It is important to document MedDRA term selection conventions, data retrieval and output strategies (including SMQs and other queries) and quality assurance procedures. Organisation-specific strategies should be consistent with the *Points to Consider* documents and should include:

- MedDRA version used for the search
- Search strategy methods (sufficiently detailed to be reproducible)
- Version update processes
- Processes for creating and maintaining customized MedDRA queries

### 2.3 – Do Not Alter MedDRA

MedDRA is a **standardised** terminology with a pre-defined term hierarchy that should not be altered. Users must not make *ad hoc* structural alterations to MedDRA, including changing the primary SOC allocation; doing so would compromise the integrity of this standard. If terms are found to be incorrectly placed in the MedDRA hierarchy, a change request should be submitted to the MSSO.

### 2.4 – Organisation-Specific Data Characteristics

Although MedDRA is a standardised terminology, different organisations have implemented it in various ways. It is important to understand organisation-specific data characteristics and implementation strategies.

Each organisation should have access to a MedDRA specialist to provide expert advice and who has the knowledge of the following database characteristics:

- Database structure (how the MedDRA hierarchy is stored and used)
- Data storage (e.g., level of term, synonym/reported term)
- Data conversion from other terminologies (if applicable)
- Coding practices over time

Example

<b>Impact of Coding Practices Over Time</b>
Consider the impact of gender-specific terms when comparing MedDRA coded data to data coded with an older terminology that may not have had corresponding gender-specific terms. If the prior terminology had only a single, gender-neutral term for “breast cancer”, consider the impact of selecting gender-specific breast cancer terms in MedDRA for current data.

- Limitations or restrictions



Example

<b>Output or Display of Multiaxial PTs</b>
Do not assume that PTs in their secondary SOC locations will be seen when searching in a specific HLT or HLGTT since the database configuration may not allow output or display by the secondary path.

- Term selection principles used
  - Selecting more than one term when coding a medical condition increases counts of terms.
  - Selecting a diagnosis term only (and not terms for signs and symptoms) reduces the counts of terms.
  - The adverse event profile resulting when both diagnosis and signs/symptoms terms are coded may appear different than when the diagnosis only is coded. Always consider the organisation's coding conventions when using or comparing data from other databases (e.g., co-developing or co-marketing partners, regulatory authorities).

## 2.5 – Characteristics of MedDRA that Impact Data Retrieval and Analysis

MedDRA's structure, rules and conventions are detailed in the MedDRA *Introductory Guide*.

Keep the following MedDRA characteristics in mind for data retrieval and presentation:

### 2.5.1 Grouping terms (HLTs and HLGTTs)

The HLT and HLGTT levels are an additional tool for data analysis and retrieval as they provide clinically relevant groupings of terms.

Example

<b>Cardiac Arrhythmias</b>
HLGTT Cardiac arrhythmias <ul style="list-style-type: none"><li>HLT Cardiac conduction disorders</li><li>HLT Rate and rhythm disorders NEC</li><li>HLT Supraventricular arrhythmias</li><li>HLT Ventricular arrhythmias and cardiac arrest</li></ul>

#### 2.5.1.1 Review terms within a grouping term

Review terms within the HLGTT or HLT of interest to be sure that all terms therein are suited for the purpose of the output.

Example

<b>Blood Pressure Terms</b>
HLT Vascular tests NEC (incl blood pressure) PT Blood pressure PT Blood pressure abnormal PT Blood pressure decreased PT Blood pressure increased
Note that terms for increased <b>and</b> decreased blood pressure are grouped under a single HLT which also includes PTs for pulmonary arterial pressure, vascular resistance, haemodynamic tests, etc.

### 2.5.2 Granularity

MedDRA PTs are more specific (“granular”) than comparable terms in other terminologies. Figure 1 illustrates how data coded to a single concept from another terminology may be coded to several PTs in MedDRA.

Related events that may have been represented by a single term in another terminology may be represented by more than one MedDRA PTs. The potential impact of this on signal detection should be kept in mind.

### 2.5.3 Multiaxiality

Multiaxiality means that a PT may exist in more than one SOC. This allows terms to be grouped in different, but medically appropriate, ways (e.g., by etiology or organ system). Each PT is assigned one primary SOC; all other SOC assignments for that PT are called “secondary”. Having a single primary SOC prevents double counting of events when outputting data from all SOCs. All possible secondary SOC assignments for any given PT may not be present in MedDRA. However, new or revised SOC assignments can be created as a result of the change request process.

#### 2.5.3.1 Primary SOC assignment rules

Primary SOC assignment rules are described in the MedDRA *Introductory Guide*. These rules affect the way terms are placed in MedDRA and determine their data display by SOC. Because these rules allow for terms related to a particular medical condition to be in more than one SOC, users should be familiar with the general structure and content of all MedDRA SOCs to be sure that data are not overlooked.

Example

Type of Disorder	Primary SOC Rule	Example	Comment
Congenital	All terms for congenital disorders have as their primary SOC assignment SOC <i>Congenital, familial and genetic disorders</i>	PT <i>Congenital absence of bile ducts</i> has a primary SOC assignment of SOC <i>Congenital, familial and genetic disorders</i> and a secondary SOC assignment of SOC <i>Hepatobiliary disorders</i>	The secondary SOC assignment for these terms is their “site of manifestation” SOC
Neoplastic	All terms for malignant and benign neoplasms (except cysts and polyps) have as their primary SOC assignment SOC <i>Neoplasms benign, malignant and unspecified (incl cysts and polyps)</i>	PT <i>Skin cancer</i> has a primary SOC assignment of SOC <i>Neoplasms benign, malignant and unspecified (incl cysts and polyps)</i> and a secondary SOC assignment of SOC <i>Skin and subcutaneous tissue disorders</i>	<b>Cyst</b> and <b>polyp</b> terms are an exception to this rule. The primary SOC assignment for cyst and polyp terms is the “site of manifestation” SOC, and the secondary SOC is SOC <i>Neoplasms benign, malignant and unspecified (incl cysts and polyps)</i>
Infectious	All terms for infectious disorders have as their primary SOC assignment SOC <i>Infections and infestations</i>	PT <i>Enterocolitis infectious</i> has a primary SOC assignment of SOC <i>Infections and infestations</i> and a secondary SOC assignment of SOC <i>Gastrointestinal disorders</i>	The secondary SOC assignment for these terms is their “site of manifestation” SOC

If a PT links to more than one of these three SOCs, the following priority is used to determine the primary SOC:

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

### 2.5.3.2 Non multiaxial SOC

Terms in the following three SOC do not have multiaxial links:

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

This is important when designing queries and other retrieval strategies because one cannot rely on multiaxiality to locate all terms of interest in MedDRA.

Example

<b>Impact of Non Multiaxial SOC on Data Queries</b>
<p>When querying a database for events or cases of thrombocytopenia, data coded to PTs in SOC <i>Blood and lymphatic system disorders</i> is a logical starting point. Additionally, data coded to terms in SOC <i>Investigations</i> – such as PT <i>Platelet count decreased</i> – and data coded to terms in SOC <i>Surgical and medical procedures</i> - such as PT <i>Platelet transfusion</i> – could also be of interest. Neither of these PTs has a link to SOC <i>Blood and lymphatic system disorders</i>.</p> <p><b>Failure to consider data coded in the non multiaxial SOC could lead to incomplete analysis of thrombocytopenia.</b></p>

As noted above, terms for test results are in SOC *Investigations* and do not have multiaxial links to terms for corresponding medical conditions. Keep this in mind when reviewing tables and data listings of MedDRA coded data.

Example

<b>Terms for Test Results in SOC Investigations</b>
<p>When querying a database for events or cases of hepatic abnormalities, data coded to PTs in SOC <i>Hepatobiliary disorders</i> is a logical starting point. Additionally, data coded to terms in SOC <i>Investigations</i> – such as PT <i>Liver function test abnormal</i> – and data coded to terms in SOC <i>Surgical and medical procedures</i> - such as PT <i>Liver transplant</i> – could also be of interest. Neither of these PTs has a link to SOC <i>Hepatobiliary disorders</i>.</p> <p><b>Failure to consider data coded in the non multiaxial SOC could lead to incomplete analysis of hepatic abnormalities.</b></p>

Figure 2 further illustrates the impact of data coded as test results vs. the corresponding medical condition.

### 2.5.3.3 Clinically related PTs

Clinically related PTs might be overlooked or not recognized as belonging together because they might be in different groupings within a single SOC or they may be located in more than one SOC. (See Section 2.5.3).

Example

<b>Similar Skin Conditions in Different Groupings</b>
HLGT Epidermal and dermal conditions
HLT Bullous conditions
PT Stevens-Johnson syndrome
PT Toxic epidermal necrolysis
HLT Exfoliative conditions
PT Dermatitis exfoliative
PT Dermatitis exfoliative generalised
PT Nikolsky's sign
PT Skin exfoliation

The frequency of a medical concept may be underestimated if the above points are not considered; this may impact interpretation of data. (See Section 3.2). MedDRA SOCs group terms by body systems, aetiologies and specialised purposes. Data may be coded to terms in SOCs that had not been anticipated by the user. Keep in mind the potential impact of multiaxiality on frequencies of the medical condition of interest.

Example

<b>Preferred Term</b>	<b>Primary SOC</b>
Post procedural haemorrhage	Injury, poisoning and procedural complications
Chest pain	General disorders and administration site conditions

## 2.6 – MedDRA Versioning

MedDRA is updated twice yearly. Version “X.0” contains both simple and complex changes; version “X.1” contains only simple changes.

Organisations should be aware of the types of MedDRA changes for their possible impact on data output.

<b>Types of MedDRA Changes</b>	
<b>Simple Changes</b>	<b>Complex Changes</b>
Add a PT (new medical concept) Move an existing PT from one HLT to another Demote a PT to LLT level Add or remove a link to an existing PT Add an LLT Move an existing LLT from one PT to another Promote an LLT to PT level Make a current LLT non-current or a non-current LLT current Changing the primary SOC allocation Changes to SMQs	Add or change multiaxial links Add new grouping terms Merge existing grouping terms Restructure a SOC

Both simple and complex changes impact retrieval and presentation strategies. Users should read the documentation provided with each MedDRA release, especially the *What's New* document. The MSSO and JMO provide tools to assist the user in comparing the changes between MedDRA versions. The Version Report (provided by the MSSO and JMO) is a spreadsheet listing all changes between the current version of MedDRA and the one previous to it; this spreadsheet is provided with each new release of MedDRA. The MSSO also provides the MedDRA Version Analysis Tool (MVAT) that facilitates identification and understanding of the impact of changes between any two MedDRA versions, including non-consecutive ones. (See Appendix, Section 6.1 of this document; also, see Section 4.1.1 of the *MedDRA Term Selection: Points to Consider* document).

Organisations should plan and document their strategy for handling MedDRA version updates. When planning or performing data retrieval and presentation, the MedDRA version used should be documented.

Keep in mind that MedDRA changes may impact previous data retrieval approaches and results, including event frequencies.

Example

<b>Impact of Version Changes – Demoted PT</b>
PT <i>Pleural haemorrhage</i> was included in a query developed using terms in MedDRA Version 16.1. If the query had been re-run on data using MedDRA Version 17.0, these events would not have been found at the PT level because PT <i>Pleural haemorrhage</i> had been demoted to an LLT and linked to PT <i>Haemothorax</i> . See Figure 3.

## Example

<b>Impact of Version Changes – Change of Primary SOC Assignment</b>
PT <i>Decreased activity</i> had a primary link to SOC <i>Psychiatric disorders</i> and a secondary link to SOC <i>General disorders and administration site conditions</i> in MedDRA Version 16.1. In Version 17.0, the primary SOC assignment was changed to SOC <i>General disorders and administration site conditions</i> and the secondary assignment to SOC <i>Psychiatric disorders</i> . In a primary SOC output of data, PT <i>Decreased activity</i> will seem to have “disappeared” from SOC <i>Psychiatric disorders</i> .

Terms used to construct queries should be in the same MedDRA version as the data being queried. An organisation’s legacy data may be coded in more than one version of MedDRA. New terms may have been included in a new query built in MedDRA Version 17.0; depending upon the organisation’s versioning method, these new terms might not be present in the older data. This could lead to search results that are incomplete.

A search built with terms of an earlier MedDRA version (e.g., used previously on a now closed study) might not identify all relevant data in an integrated safety summary (ISS) containing data coded in a later version of MedDRA. Queries stored in an organisation’s system should be updated to the appropriate version of MedDRA before using them on new data.

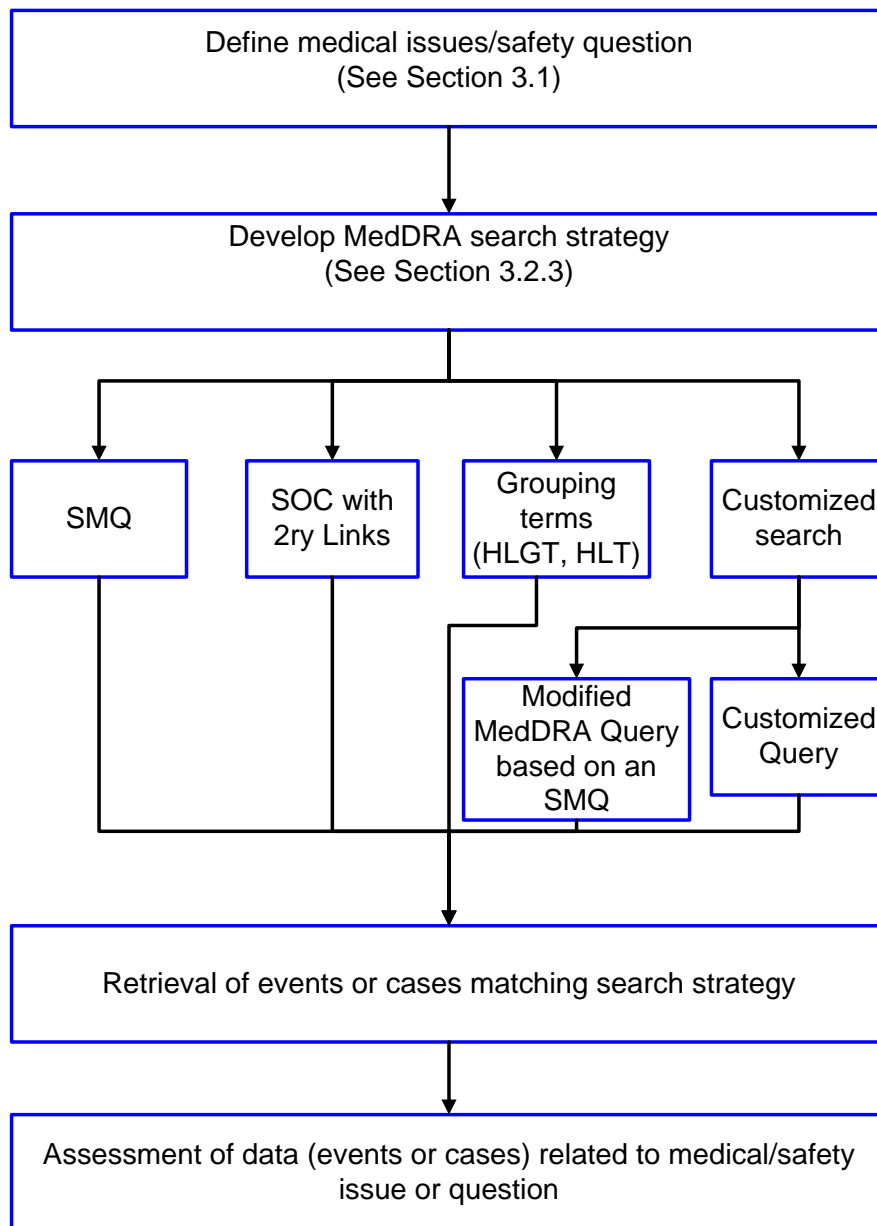
Advice on how an organisation should handle new MedDRA versions is not within the scope of this document. (See *MedDRA Term Selection: Points to Consider*, Appendix 4.1). Some databases may contain data of multiple studies coded in different versions of MedDRA. This may impact aggregation of those data (e.g., in an ISS). Refer to MSSO documents on versioning options for clinical trial and post-marketing data for more information. (See Appendix, Section 6.1).

## **Section 3 – GENERAL QUERIES AND RETRIEVAL**

### **3.1 – General Principles**

Data retrieval is performed for summary and analysis of clinical trial data, pharmacovigilance, medical information questions and for a number of other purposes. The search strategies, methods and tools used to retrieve data might differ based on the intended use of the output.

A general approach for data retrieval is outlined in the chart below.



Prior to data retrieval, there may be known or potential safety issues that need detailed investigation. Information from pre-clinical studies, clinical trials post-marketing surveillance, class effects of similar products, and regulatory queries may identify areas of possible focus; these may affect the strategy for aggregating search terms, the methodology, and the way data are displayed.

Be aware of database characteristics, organisation-specific data entry conventions, data sources, the size of the database, and the version of MedDRA used for coding all data. Archived searches may be available to the user, especially those used in pharmacovigilance; these may be suitable for use if updated.



When presenting adverse event data, it is important to display and to group related events (i.e., events that represent the same condition of interest) so that the true occurrence rate of an event is not obscured. **Search strategies should be documented.** The search output alone may not suffice for data assessment (e.g., frequency of a condition). Search results should be evaluated against the question originally posed.

Sorting related events into categories can be challenging. A search that is too narrowly focused might exclude events of potential relevance; a search that is too broad might make it difficult to identify a trend or signal. Careful interpretation is required when grouping terms that correspond to a potential event or medical condition for analysis (whether a syndrome or not). The purpose is to identify trends that may require further analysis, including review of individual cases. For complex queries, create a data analysis plan including a definition of the medical condition of interest. An interdisciplinary discussion might be helpful to identify the most suitable methods and tools relevant to the query.

These principles may apply to the types of searches listed in the table below:

Example

<b>Types of Searches – Application of General Principles</b>
Safety profile overview in a summary report, Periodic Safety Update Report (PSUR), ISS, etc.
Comparing frequencies of ARs/AEs reporting rates for spontaneous reports or incidence for studies)
Analysis of a specific safety concern
Identifying patient subpopulations at risk (search of medical history)

### **3.1.1 Graphical displays**

Graphical displays can be useful especially with large datasets. Such displays allow quick visual representation of potential signals. Organisations are encouraged to use graphs for data display. Histograms, bar charts, and pie charts can be useful as can more complex, statistically-derived displays (e.g., data mining algorithms). Examples of these types of displays are in the Appendix, Section 6.3.

### **3.1.2 Patient subpopulations**

Paediatric and gender-specific adverse event terms lists – available on the MedDRA and JMO websites – may aid in data retrieval for these subpopulations (See Appendix, Section 6.1). However, it is necessary to refer to individual database fields for demographics.

### 3.2 – Overall Presentation of Safety Profiles

The aims of an overall safety profile presentation are to:

- Highlight distribution of ARs/AEs
- Identify areas for in depth analysis

Present the data in a way that allows for easy recognition of patterns of terms potentially related to the relevant medical conditions. There are various ways to do this ranging from a full listing of terms to sophisticated statistical approaches such as data mining techniques (for reference, see ICH E2E: Pharmacovigilance Planning Document; listed in the Appendix, Section 6.1).

Historically, the standard approach has been to display data by Body System (or System Organ Class) and Preferred Term corresponding to SOCs and PTs in MedDRA. Due to MedDRA's unique characteristics (multiaxiality, granularity), this PT-SOC approach may need to be augmented with other types of data outputs (e.g., secondary SOC output, display by grouping terms [HLTs, HLGTS], etc.), depending on the reason for the output. For example, if a number of reports describe a similar medical condition, they could be represented by:

- Many different PTs (dilution of signal)
- Different grouping terms
- Different SOCs

SOCs where the user would not intuitively expect them (e.g., SOC *General disorders and administration site conditions*, SOC *Pregnancy, puerperium and perinatal conditions*, SOC *Injury, poisoning and procedural complications*, SOC *Infections and infestations*). See examples in the table below

Example

<b>PTs with Primary SOC <i>General disorders and administration site conditions</i> and Secondary SOC <i>Cardiac disorders</i></b>
PT Chest discomfort
PT Chest pain
PT Oedema peripheral
PT Sudden death
PT Localised oedema
PT Oedema due to cardiac disease
PT Peripheral oedema neonatal
PT Cardiac death

### 3.2.1 Overview by primary System Organ Class

This overview is recommended as a first step in data retrieval and for planning of further analysis.

Display of all data ensures that all events will be seen and may be useful to identify data clusters by SOC. If the hierarchy is also displayed, clusters may occur at the HLG or HLT levels. For a small dataset, this display by primary SOC may be all that is necessary.

- Objectives:
  - Include all events (none are omitted)
  - Display all data in the entire MedDRA hierarchy
- Method:

The primary SOC view including HLGs, HLTs and PTs can be used for standard tables (clinical trials and post-marketing data) and for cumulative summaries (post-marketing data). Line listings (both clinical and post-marketing data) can also be displayed by primary SOC and PT. Depending on the reason for the output, it might be beneficial to use the primary SOC and PT display; for large datasets, display by SOC **and** by grouping terms (HLGs and HLTs) may be preferable. Figure 4 is an example of such an output.

The Internationally Agreed Order of SOCs was developed for consistency irrespective of language or alphabet (see Figure 5). The SOC order was based upon the relative importance of each SOC in AR/AE reports. Use of the Internationally Agreed Order may be applicable to certain regulatory functions, e.g., the SPC guideline (see the MedDRA *Introductory Guide* and MedDRA ASCII files). Organisations that share data should agree on the order of SOCs when preparing data for presentation.

Data displays in tables or in graphical presentations may facilitate understanding by the viewer. Figures 6, 7 and 8 are examples of such displays.

Figures 9a and 9b display data for one compound in two patient populations. Within each patient population, the reports are split by SOC and by reporter. The upper bar of each pair represents numbers of reports from consumers (blue), and the lower bar represents reports from health care professionals (red). If further detail is needed, adverse events can be displayed by PT with decreasing frequency.

In depth analysis requires medical expertise to define terms that should be aggregated.

- Benefits:
  - Provides an overview of data distribution; helps identify areas of special interest that may need in depth analysis
  - Grouping terms aggregate related PTs, facilitating identification of medical conditions of interest
  - A PT will be displayed only once, preventing over-counting of terms
  - A primary SOC overview may be the only form of data display necessary for a small dataset
- Limitations:
  - Because it is based on a PT-to-primary SOC assignment, there may be incomplete groupings of terms for a medical condition or syndrome as such terms may be distributed among different SOCs
  - Events may not be found where the user expects them due to MedDRA placement rules
  - Potential for a lengthy data output when applied to large datasets

### **3.2.2 Overall presentations of small datasets**

When the safety profile consists of a small list of PTs (e.g., early in clinical development), a display of these PTs may be adequate. Figure 10 is an example of this.

### **3.2.3 Focused searches**

Focused searches may be useful for further investigation of medical concepts of interest. For example, a focused search may be used to determine the number of cases or events of interest in response to a regulatory query.

In certain situations, such as those listed below (note that this list is not all-inclusive), users may wish to design a specific search in addition to the Overview by Primary System Organ Class.(See Section 3.2.1).

- Further examination of clusters seen in Primary SOC output
- Previously identified safety concerns (e.g., known class effects, results from toxicology and animal studies, etc.)
- Monitoring events of special interest
- Responding to regulatory queries

Below are listed options for focused search approaches. The order of applying these approaches may depend on resources, expertise, systems or other factors.

### 3.2.3.1 Focused searches by secondary SOC assignments

This focused search augments the Overview by Primary System Organ Class (see Section 3.2.1) by addressing secondary SOC assignments, thus providing a more comprehensive view of the data and taking advantage of MedDRA's multiaxiality.

➤ Method:

The method used for a focused search by secondary SOC assignment may depend on the database characteristics of the organisation.

Options include:

- Query of the SOC, HLG and HLT levels to include both the primary and secondary SOC assignments in the display
- Output PTs in their secondary SOC locations programmatically (See Figure 11)
- If the database does not allow automated output by secondary SOC, then perform the query using available processes (e.g., programming a list of all individual PTs in the primary and secondary SOC locations)

Example

<b>Programming a List of PTs in Primary and Secondary SOC Locations</b>
SOC Eye disorders HLG Vision disorders HLT Visual pathway disorders PT Chiasma syndrome <b>PT Optic nerve disorder (primary SOC location)</b> <b>PT Optic neuropathy (primary SOC location)</b> <b>PT Toxic optic neuropathy (primary SOC location)</b> PT Visual cortex atrophy PT Visual pathway disorder
<b>3 of 6 PTs are primary to SOC Nervous system disorders</b>

➤ Benefits:

Multiaxial links enhance the utility of the grouping terms. This method overcomes the primary SOC limitations as described under Section 3.2.1.

➤ Limitations:

- Still displays only terms that are represented in one SOC or HLG/HLT which may not include all terms related to a medical condition
- This method of display of PTs by both primary and secondary SOC assignments could lead to double counting of cases/events

## **Section 4 – STANDARDISED MedDRA QUERIES**

### **4.1 – Introduction**

Standardised MedDRA Queries (SMQs) were created to standardise identification and retrieval of safety data.

SMQs are a joint effort of the Council for International Organizations of Medical Sciences (CIOMS) and ICH (including MSSO and JMO) representing both industry and regulatory authorities. An SMQ is a grouping of terms from one or more SOCs that relate to a defined medical condition or area of interest. The terms included relate to signs, symptoms, diagnoses, syndromes, physical findings, laboratory and other physiologic test data, etc. that are associated with the medical condition or area of interest.

Users should carefully read the *Introductory Guide for Standardised MedDRA Queries (SMQs)* before applying an SMQ to fully understand the scope of the SMQ and to properly apply search options such as algorithms and weightings.

### **4.2 – SMQ Benefits**

As with all MedDRA-based queries, users of SMQs should be aware of several factors that may influence data retrieval including database characteristics, data conversion processes, coding conventions, and MedDRA versioning. For more details, see Section 3.1.

➤ SMQ benefits include:

- Application across multiple therapeutic areas
- Validated reusable search logic
- Standardised communication of safety information

- Consistent data retrieval
- Maintenance by MSSO and JMO

### **4.3 – SMQ Limitations**

- SMQs do not cover all medical topics or safety issues
- SMQs evolve and undergo further refinement even though they have been tested during development

### **4.4 – SMQ Modifications and Organisation-Constructed Queries**

If any modifications are made to term content or structure of an SMQ, it can no longer be called an “SMQ” but it should instead be referred to as a “modified MedDRA query based on an SMQ”. See Section 5.1 for further details on SMQ modification.

**Under no circumstances should a query constructed for the specific need of an organisation be called an “SMQ” by its originator.** This is to ensure that there is no confusion with the ICH-endorsed SMQs applied by other MedDRA users. Any alternate name for the organisation-constructed query is acceptable as long as it could not be potentially confused with an ICH-endorsed SMQ.

### **4.5 – SMQs and MedDRA Version Changes**

Each SMQ relates to a specific MedDRA version. SMQs are part of each new MedDRA release, are maintained by MSSO and JMO, and correspond to the terms present in that version of MedDRA. The SMQ version should always correspond to the MedDRA version of the data being searched.

As with all searches of MedDRA-based data, it is important to document the MedDRA and SMQ versions used.

Changes to SMQs that can occur with each MedDRA version include (but are not limited to) the following:

- Addition of PTs
- Inactivation of a PT (i.e., effectively “removing” a PT from an SMQ)
- Change of term scope (e.g., a narrow term becomes a broad term)
- Restructuring of an SMQ (e.g., change in the hierarchical position of an SMQ)

- Creation of a new SMQ

For a full description of the types of changes that can occur to SMQs, please refer to the MedDRA “Change Request Information” document. (See Appendix, Section 6.1). Changes introduced with each new version are documented in the “What’s New” document for each MedDRA version. (The cumulative changes are contained within the ASCII files in the fields called “Term\_addition\_version” and “Term\_last\_modified\_version”).

The MedDRA version of the SMQ and the coded data being searched should be the same because mismatches could produce unexpected results. For example, if an SMQ from an older version of MedDRA is applied to data coded in a more recent version, data coded to terms that are not present in the older SMQ would not be retrieved.

Example

<b>Consequence of Version Mismatch of Coded Data and SMQ</b>
PT <i>Splenic artery thrombosis</i> was added to SMQ <i>Embolic and thrombotic events, arterial</i> in MedDRA Version 17.0. Using Version 16.1 of this SMQ – which does not contain this PT – would fail to identify cases coded to this term in a database using MedDRA Version 17.0.

#### **4.6 – SMQs – Impact of MedDRA Legacy Data Conversion**

The conversion method for data originally coded in another terminology (e.g., COSTART) also impacts the application and output of SMQs. See Section 2.1.2, *Impact of data conversion method*.

#### **4.7 – SMQ Change Requests**

Users are encouraged to submit Change Requests to MSSO and JMO to improve the utility of SMQs. A justification (and possibly testing data) for a submitted Change Request must be provided. The MSSO may require more time to evaluate SMQ Change requests than regular MedDRA Change Requests. Before submitting an SMQ Change Request, users should review the SMQ documentation for inclusion and exclusion criteria of the SMQ.

#### **4.8 – SMQ Technical Tools**

The MSSO browsers (both the desktop and Web-based browsers) allow for searching and viewing the contents of SMQs and they include additional details such as the SMQ description (definition) and development notes. An Excel spreadsheet containing the terms in each production SMQ is available from MSSO and JMO. (See Appendix, Section 6.1). This spreadsheet allows a user to transfer SMQ terms to query tools. File specifications related to SMQs are found



in the “MedDRA Distribution File Format Document” supplied with each MedDRA version.

The MedDRA website has a list of some system tools that provide technical support for SMQs. (See Appendix, Section 6.1).

## 4.9 – SMQ Applications

SMQs were developed to address the high granularity and unique features of MedDRA and to maximise the likelihood that all terms related to a specific medical condition of interest are identified.

The user should first review the list of available SMQs to determine which of them may be applicable to the question being asked. If an SMQ seems applicable, the user should check the documentation in the SMQ Introductory Guide to understand the purpose and definition of the SMQ. The user may also wish to review the term contents of the SMQ.

Following application of the selected SMQ on coded data, search results (i.e., retrieved data) should then be evaluated against the question originally posed. The search output alone may not be sufficient for data assessment (e.g., frequency of a condition). Define and document criteria for case evaluation.

Generally, more cases/events will be retrieved than will eventually be subjected to analysis due to “noise”. This is a more significant consideration for “broad” searches but in principle also applies to “narrow” searches. (See Section 4.10.1).

### 4.9.1 Clinical trials

SMQs may be applied in the clinical trial setting – especially for aggregate data – where the safety profile has yet to be fully established. In this instance, most (if not all) available SMQs may be used, possibly on a routine basis.

Alternatively, a user can apply an SMQ (or SMQs) that relates to a previously identified area of interest (e.g., from pre-clinical data or class effect) for further evaluation.

Example

<b>Targeted Safety Study</b>
When developing a data analysis plan for a targeted safety study, consider using the narrow terms of an SMQ to aggregate events of interest.

## 4.9.2 Post-marketing

### 4.9.2.1 Focused searches

A specific SMQ or a selection of SMQs may be used to retrieve relevant cases for subsequent medical review.

Example

<b>Emerging Safety Signal</b>
A company suspects an emerging signal of pancreatitis for a new HIV product. SMQ <i>Acute pancreatitis</i> can be applied to the data.

### 4.9.2.2 Signal detection

The entire set of SMQs may be used on the database for signal detection. The user may wish to use the narrow terms or more specific levels of hierarchical SMQs (i.e., a sub-search SMQ) to minimise dilution of the signal.

### 4.9.2.3 Single case alert

SMQs may be used to create a “watch list” (e.g., an automated notification system) to alert the user of incoming cases needing urgent review.

Example

<b>Single Case Alert</b>
A medical issue of interest needs to be communicated to a regulatory authority as part of an agreed risk management plan. The SMQ narrow search or more specific levels of a hierarchical SMQ may be applied to identify potential cases of interest.

### 4.9.2.4 Periodic reporting

SMQs may help aggregate relevant cases for ongoing review of specific safety issues in periodic safety reports. SMQs may also be used for other routine reviews of aggregate data (e.g., reports of lack of efficacy) in the context of a periodic report.

## 4.10 – SMQ Search Options

Some SMQs have options that may be used to refine a particular search. The most common option is use of narrow and broad search terms. By definition, a broad search includes both narrow and broad terms.

Some SMQs are hierarchical (i.e., contain one or more sub-searches). Other SMQs use algorithms, and in one case (SMQ *Systemic lupus erythematosus*),

weightings are assigned to particular terms for signs, symptoms and laboratory results to help identify cases.

#### 4.10.1 Narrow and broad searches

Most SMQs have narrow and broad PTs. The narrow PTs have a greater likelihood of identifying only events of interest (high specificity) while the broad terms are intended to identify additional possible events (high sensitivity). Some events retrieved by the broad search terms may, upon further review, not relate to the condition of interest. The user can select the scope of the search (narrow or broad) that is most applicable to the question being asked. Figure 12 is an example of output of narrow and broad searches.

When a compound is in early phase development or has only recently been marketed, it may be advisable to use the broad search.

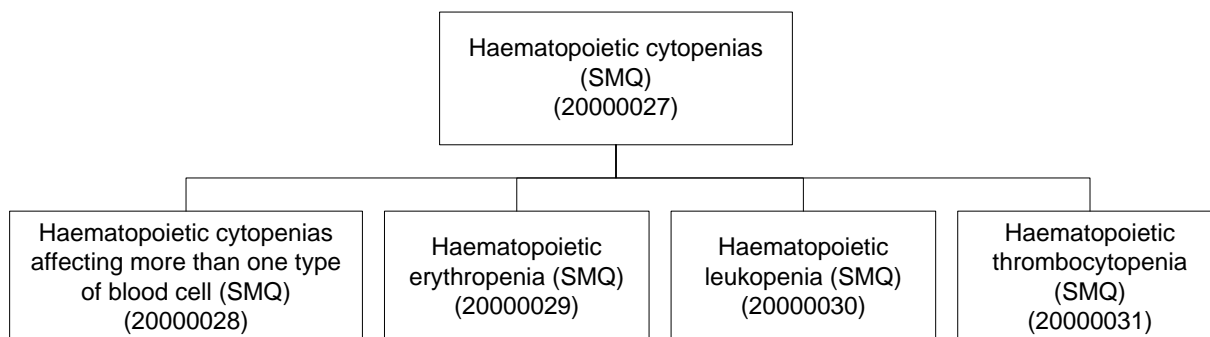
Example

Use of Broad Search
If evaluating an emerging signal of lactic acidosis using SMQ <i>Lactic acidosis</i> , narrow terms may be applied to identify events where the specific diagnosis has been reported; however, events of reported signs and symptoms would not be retrieved. If there is additional need to find cases where no specific diagnosis (but mainly signs and symptoms) have been reported, then a broad search (i.e., narrow + broad search terms) should be applied.

#### 4.10.2 Hierarchical SMQs

Several SMQs have a hierarchical structure (one or more levels of sub-searches of increasing specificity). The user can select the search that is most applicable to the question being asked or a combination of sub-search SMQs as needed.

The SMQ Introductory Guide has explanatory notes on the appropriate use of each hierarchical SMQ. An example of a hierarchical SMQ is illustrated below (SMQ *Haematopoietic cytopenias*).



## Example

Use of SMQ Hierarchy
The medical condition of interest is thrombocytopenia. SMQ <i>Haematopoietic cytopenias</i> may be too inclusive because sub-searches for decreases of other hematopoietic cell lines (e.g., SMQ <i>Haematopoietic leukopenia</i> ) are included. A user may wish to select only the sub-search SMQ <i>Haematopoietic thrombocytopenia</i> in this instance.

### 4.10.3 Algorithmic SMQs

An algorithm provides for a combination of terms which – if retrieved in a single case – are more likely to identify a case of interest than isolated broad search terms (see table below). The broad terms of algorithmic SMQs are subdivided into **categories** that could be groupings of organ-specific signs or symptoms, laboratory terms, etc. (Note: the broad search categories are labeled B, C, D, etc.). Using an algorithm may reduce the amount of “noise” (i.e., non-relevant cases).

Using an algorithmic SMQ without applying the algorithm (i.e., simply applying the narrow and broad searches) will yield different results from those obtained using the algorithm.

## Example

Algorithmic SMQ (SMQ <i>Anaphylactic reaction</i> )*		
Category B – Upper airway/Respiratory	Category C – Angioedema/Urticaria, etc.	Category D – Cardiovascular/Hypotension
Acute respiratory failure	Allergic oedema	Blood pressure decreased
Asthma	Angioedema	Blood pressure diastolic decreased
Bronchial oedema	Erythema	Blood pressure systolic decreased
Algorithm: <ul style="list-style-type: none"><li>• Case = A (Narrow terms – not included in the table)</li><li>• Or term from Category B <b>and</b> term from Category C</li><li>• Or term from <b>either</b> Category B or Category C <b>plus</b> term from Category D</li></ul>		

\* Not all terms in these categories are listed in the table

SMQ *Systemic lupus erythematosus* is an algorithmic SMQ with assigned weights for its included PTs (e.g., PT *Pleural effusion* = 3); a total weighted score greater than 6 suggests a case of interest.

Users should not assume that all software tools support algorithmic SMQs.

#### 4.11 – SMQ and MedDRA Grouping Terms

Data retrieved using MedDRA grouping terms (HLGTs, HLTs) may differ from those retrieved using a related SMQ.

Example

Comparison – SMQ and Grouping Terms
Cardiac arrhythmia is a suspected issue (e.g., by review of a primary SOC output of all data). If events retrieved by using HLGT <i>Cardiac arrhythmias</i> are compared to those retrieved by SMQ <i>Cardiac arrhythmias</i> , more events may be retrieved by the SMQ because it includes additional terms from other SOCs such as SOC <i>Investigations</i> .

### Section 5 – CUSTOMISED SEARCHES

MedDRA allows for a variety of searching options as described above. However, there will be situations when a customised search is needed.

#### 5.1 – Modified MedDRA Query Based on an SMQ

Do not modify the term content or structure of an SMQ unless there is a compelling reason to do so since altering it in any way makes it non-standard. (See Section 4.4).

If an SMQ is modified in any way, it should be referred to as a “modified MedDRA query based on an SMQ”. All modifications to the original SMQ should be documented.

If a modified MedDRA query based on an SMQ is to be used on an ongoing basis, version updates and maintenance of the query are the responsibility of the organisation that created it.

## Example

Modified MedDRA Queries based on SMQs	
Additional PTs are needed	A product is being investigated for a possible safety signal of dementia, and the user wishes to use SMQ <i>Dementia</i> . For this particular product, PT <i>Disturbance in attention</i> may be needed.
Exclusion of PTs	An antipsychotic product is being investigated for potential QT prolongation and also has a well-described association with hypotension and fainting. When using SMQ <i>Torsade de pointes/QT prolongation</i> (broad search), the user may wish to exclude PT <i>Syncope</i> to prevent excess “noise” in data retrieval.
Changing the scope (narrow or broad) of an SMQ term	A product is being investigated for the potential for hyperglycaemia and diabetes mellitus. SMQ <i>Hyperglycaemia/new onset diabetes mellitus</i> has PT <i>Increased insulin requirement</i> as a <b>broad</b> search term. For this query, it may be useful to include PT <i>Increased insulin requirement</i> in the <b>narrow</b> search.

## 5.2 – Customised Queries

Consider these points when constructing a customised query for MedDRA-coded data:

- Those responsible for constructing a customised query should:
  - Have medical knowledge
  - Know the structure and characteristics of MedDRA (e.g., hierarchy, multiaxiality) and the general content of MedDRA groupings (SOCs, HLGs, and HLTs)
  - Understand the characteristics and structure of the data
- The specificity of the search should be defined.
- Initial focus should be on SOCs related to the condition of interest. For example, a customised search for a renal condition should start with SOC *Renal and urinary disorders*.
- The non multiaxial SOCs (SOC *Investigations*, SOC *Surgical and medical procedures* and SOC *Social circumstances*) should always be reviewed. Also, it may be useful to review terms in other SOCs that are not organ systems (e.g., SOC *General disorders and administration site conditions*,

*SOC Injury, poisoning and procedural complications and SOC Pregnancy, puerperium and perinatal conditions).*

- It may be useful to identify relevant query terms by the following approaches:
  - A “bottom-up” survey of MedDRA (terms at the LLT and PT levels initially)
  - A “top-down” survey of MedDRA (starting at the SOC level and drilling down through the hierarchy)
- Consider looking at secondary links for multiaxial terms since additional relevant query terms could be found. For example, PT *Dyspnoea* can be found with other respiratory symptoms PTs in its primary SOC *Respiratory, thoracic and mediastinal disorders*, and it can also be found with related cardiac symptoms in its secondary SOC *Cardiac disorders*.
- Include grouping terms (HLGTs, HLTs) when possible (remembering the caveats described in Section 2.5.1).
- In general, queries should be built on PTs and grouping terms. Unless very specific concepts (e.g., bacterial species) are needed, avoid using LLTs to build queries.
- Consider saving the customised query for future use; maintenance is necessary for MedDRA version changes.
- A customised query that may be useful to other MedDRA users can be submitted to the MSSO as a Change request for possible development as an SMQ.

## Section 6 – APPENDIX

### 6.1 – Links and References

The following documents and tools can be found on the MedDRA website: ([www.meddra.org](http://www.meddra.org)):

- MedDRA Term Selection: Points to Consider document (also available on the JMO website: [www.pmrj.jp/jmo/php/indexe.php](http://www.pmrj.jp/jmo/php/indexe.php))
- MedDRA Introductory Guide
- Introductory Guide for Standardised MedDRA Queries (SMQs)
- Pediatric and Gender Adverse Event Term Lists
- MedDRA Change Request Information document
- MedDRA Web-based Browser
- MedDRA Desktop Browser
- MedDRA Version Report (lists all changes in new version) \*
- MedDRA Version Analysis Tool (compares any two versions) \*
- Production SMQ spreadsheet\*
- List of system tools that support SMQs

\* Requires user ID and password to access

The following document can be found on the ICH website ([www.ich.org](http://www.ich.org)):

- ICH E2E: Pharmacovigilance Planning



## 6.2 – Membership of the ICH Points to Consider Working Group

### 6.2.1 Current members of the ICH Points to Consider Working Group

Affiliation	Member
Commission of the European Communities	Sarah Vaughan
	Maria Luisa Casini
European Federation of Pharmaceutical Industries and Associations	Hilary Vass*
	Christina Winter†
Health Canada	Lynn Macdonald
Japanese Maintenance Organization	Yutaka Nagao
	Kazuyuki Sekiguchi
	Reiji Tezuka
Japan Pharmaceutical Manufacturers Association	Yo Tanaka
MedDRA MSSO	Judy Harrison
Ministry of Health, Labour and Welfare/Pharmaceuticals and Medical Devices Agency	Sonoko Ishihara
	Makiko Isozaki
	Yuuhei Fukuta
Pharmaceutical Research and Manufacturers of America	Anna-Lisa Kleckner
	JoAnn Medbery
US Food and Drug Administration	Sonja Brajovic#
	Christopher Breder

\* Current Rapporteur

# Regulatory Chair

† Former Rapporteur

## 6.2.2 Former members of the ICH Points to Consider Working Group

Affiliation	Member
Commission of the European Communities	Dolores Montero
	Carmen Kreft-Jais
	Morell David
European Federation of Pharmaceutical Industries and Associations	Barry Hammond <sup>†</sup> ; Reinhard Fescharek <sup>†</sup>
Health Canada	Alison Bennett, Heather Morrison; Michelle Séguin; Heather Sutcliffe; Bill Wilson
Japanese Maintenance Organization	Osamu Handa; Akemi Ishikawa; Yasuo Sakurai; Yuki Tada
Japan Pharmaceutical Manufacturers Association	Takayoshi Ichikawa; Akemi Ishikawa; Satoru Mori; Yasuo Sakurai; Kunikazu Yokoi
MedDRA MSSO	JoAnn Medbery; Patricia Mozzicato
Ministry of Health, Labour and Welfare/Pharmaceuticals and Medical Devices Agency	Tamaki Fushimi; Wakako Horiki; Kazuhiro Kemmotsu; Tatsuo Kishi; Chie Kojima; Emiko Kondo; Hideyuki Kondou; Kemji Kuramochi; Tetsuya Kusakabe; Kaori Nomura; Izumi Oba; Shinichi Okamura; Yoshihiko Sano; Nogusa Takahara; Kenichi Tamiya; Daisuke Tanaka; Shinichi Watanabe; Takashi Yasukawa; Go Yamamoto; Manabu Yamamoto; Nobuhiro Yamamoto
Pharmaceutical Research and Manufacturers of America	David Goldsmith; Sidney Kahn; Susan M. Lorenski; Margaret M. Westland <sup>†</sup>
US Food and Drug Administration	Miles Braun; Andrea Feight; John (Jake) Kelsey <sup>†</sup> ; Brad Leissa; Toni Piazza-Hepp

<sup>†</sup> Former Rapporteur

### 6.3 – Figures

OTHER TERMINOLOGY PREFERRED TERMS	No. OF EVENTS	MEDDRA VERSION 17.0 PREFERRED TERMS	No. OF EVENTS
Infection	15	Upper respiratory tract infection	7
		Nasopharyngitis	2
		Infection	1
		Lower respiratory tract infection	4
		Skin infection	1
Abdominal pain	9	Abdominal pain	4
		Abdominal pain upper	3
		Abdominal tenderness	2
Accidental injury	4	Injury	1
		Skin laceration	1
		Ligament sprain	1
		Back injury	1

Figure 1 – How data coded to a single concept from another terminology may be expressed by several PTs in MedDRA

Reported Event (% subjects)	OTHER TERMINOLOGY		MedDRA Version 17.0	
	Coded Term (% subjects)	Body System/SOC (% subjects)	PT (% subjects)	SOC (% subjects)
Hyperglycaemia (4.1)	Hyperglycaemia (10.5)	Metabolism & nutritional disorders (10.5)	Hyperglycaemia (4.1)	Metabolism and nutrition disorders (4.1)
Increased blood sugar (2.7)				
Glucose increased (2.2)			Blood glucose increased (6.4)	Investigations (6.4)
Blood glucose high (1.0)				
Increasing glucoses (0.5)				

Figure 2 – Multiple MedDRA terms may be used to code similar medical conditions included in a “disorder SOC”; associated laboratory findings are in SOC Investigations

Preferred Terms	Events/Cases		Comment
	Version 16.1	Version 17.0	
Pleural haemorrhage	15	0 (no longer a PT)	In MedDRA Version 16.1, <i>Pleural haemorrhage</i> was a PT and in Version 17.0 it was demoted to an LLT under PT <i>Haemothorax</i>
Haemothorax	5	20	

Figure 3 – Impact of MedDRA version changes – demotion of a PT

SOC (System Organ Class)		Number of all reactions*	Number of reactions* where outcome of report was fatal
HLGT	HLT   PT		
<b>Nervous system disorders</b>			
<b>Mental impairment disorders</b>			
Mental impairment (excl dementia and memory loss)	Disturbance in attention	1	0
<b>Movement disorders (incl parkinsonism)</b>			
Dyskinesias and movement disorders NEC	Psychomotor hyperactivity	2	0
Tremor (excl congenital)	Tremor	3	0
<b>Neurological disorders NEC</b>			
Disturbances in consciousness NEC	Somnolence	1	0
Neurological signs and symptoms NEC	Dizziness	1	0
<b>Seizures (incl subtypes)</b>			
Seizures and seizure disorders NEC	Convulsion	2	0
<b>Nervous system disorders SOC Total</b>		<b>10</b>	<b>0</b>
<b>Psychiatric disorders</b>			
<b>Anxiety disorders and symptoms</b>			
Anxiety symptoms	Activation syndrome	1	0
	Agitation	2	0
	Anxiety	2	0
	Stress	1	0
<b>Depressed mood disorders and disturbances</b>			
Depressive disorders	Depression	1	0
<b>Disturbances in thinking and perception</b>			
Thinking disturbances	Thinking abnormal	1	0
<b>Schizophrenia and other psychotic disorders</b>			
Psychotic disorder NEC	Psychotic disorder	1	0
<b>Sleep disorders and disturbances</b>			
Disturbances in initiating and maintaining sleep	Insomnia	1	0
<b>Psychiatric disorders SOC Total</b>		<b>10</b>	<b>0</b>

Figure 4 – Primary SOC output listing – example. Note that some PTs are multi-axial, however, this figure shows only the primary SOC assignments

<b>MedDRA Version 17.0 English Alphabetical Order</b>	<b>MedDRA Version 17.0 Internationally Agreed Order</b>
Blood and lymphatic system disorders	Infections and infestations
Cardiac disorders	Neoplasms benign, malignant and unspecified (incl cysts and polyps)
Congenital, familial and genetic disorders	Blood and lymphatic system disorders
Ear and labyrinth disorders	Immune system disorders
Endocrine disorders	Endocrine disorders
Eye disorders	Metabolism and nutrition disorders
Gastrointestinal disorders	Psychiatric disorders
General disorders and administration site conditions	Nervous system disorders
Hepatobiliary disorders	Eye disorders
Immune system disorders	Ear and labyrinth disorders
Infections and infestations	Cardiac disorders
Injury, poisoning and procedural complications	Vascular disorders
Investigations	Respiratory, thoracic and mediastinal disorders
Metabolism and nutrition disorders	Gastrointestinal disorders
Musculoskeletal and connective tissue disorders	Hepatobiliary disorders
Neoplasms benign, malignant and unspecified (incl cysts and polyps)	Skin and subcutaneous tissue disorders
Nervous system disorders	Musculoskeletal and connective tissue disorders
Pregnancy, puerperium and perinatal conditions	Renal and urinary disorders
Psychiatric disorders	Pregnancy, puerperium and perinatal conditions
Renal and urinary disorders	Reproductive system and breast disorders
Reproductive system and breast disorders	Congenital, familial and genetic disorders
Respiratory, thoracic and mediastinal disorders	General disorders and administration site conditions
Skin and subcutaneous tissue disorders	Investigations
Social circumstances	Injury, poisoning and procedural complications
Surgical and medical procedures	Surgical and medical procedures
Vascular disorders	Social circumstances

*Figure 5 – The alphabetical SOC order (in English) and the Internationally Agreed Order of SOCs*

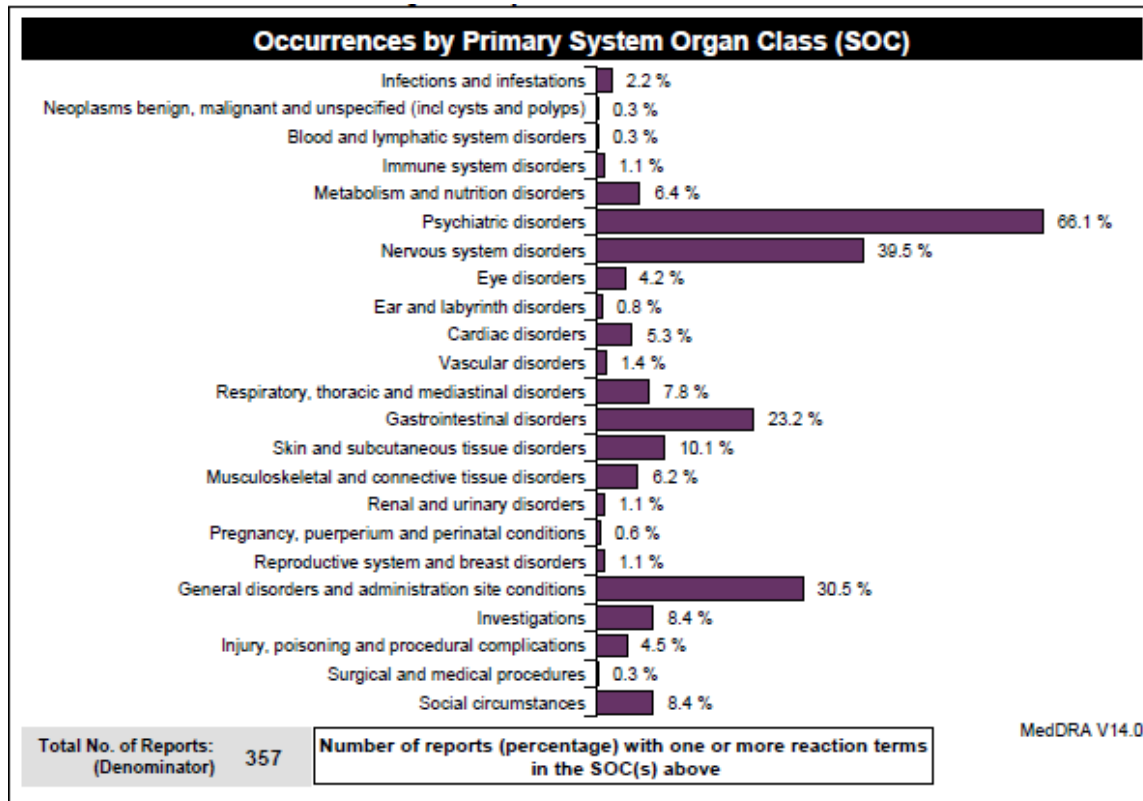


Figure 6 – Example of a graphical display (frequency by primary SOC)

Figure 2  
Relative frequency of events per primary [1] and per secondary [2] SOC

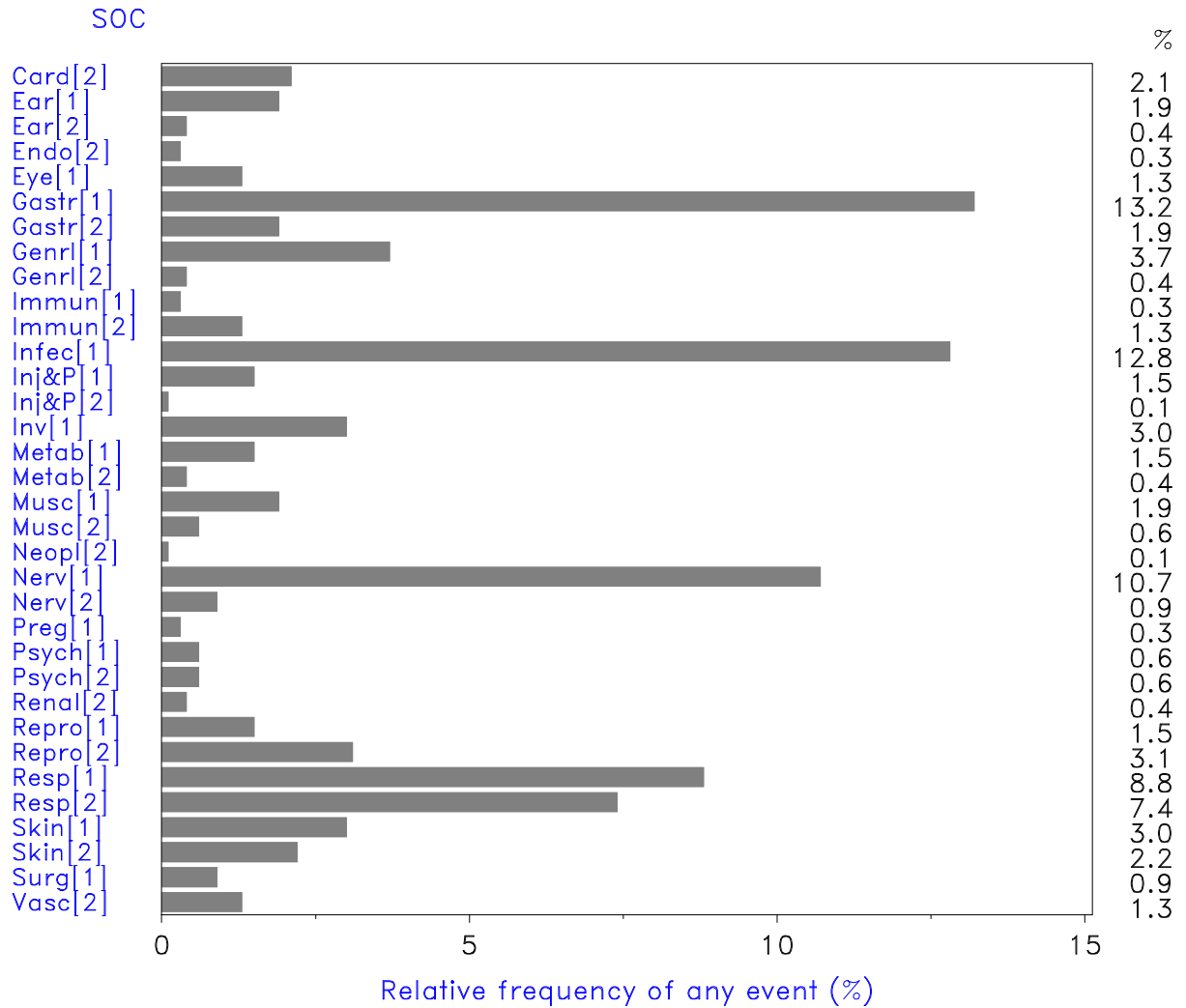


Figure 7 – Example of a graphical display (frequency by primary and secondary SOC)

System Organ Class	Number of All Reactions*	Reactions* (% of total)	Number of Reactions* where outcome of AR report was fatal
Gastrointestinal disorders	1	1.92 %	0
General disorders and administration site conditions	10	19.23 %	0
Hepatobiliary disorders	2	3.85 %	0
Immune system disorders	1	1.92 %	0
Infections and infestations	1	1.92 %	0
Investigations	7	13.46 %	0
Metabolism and nutrition disorders	1	1.92 %	0
Musculoskeletal and connective tissue disorders	1	1.92 %	0
Nervous system disorders	10	19.23 %	0
Psychiatric disorders	10	19.23 %	0
Renal and urinary disorders	2	3.85 %	0
Respiratory, thoracic and mediastinal disorders	2	3.85 %	0
Skin and subcutaneous tissue disorders	4	7.69 %	0
<b>Total Number of Reactions:</b>	<b>52</b>	<b>100.00 %</b>	<b>0</b>

Figure 8 – Example of a tabular display (frequency by primary SOC)

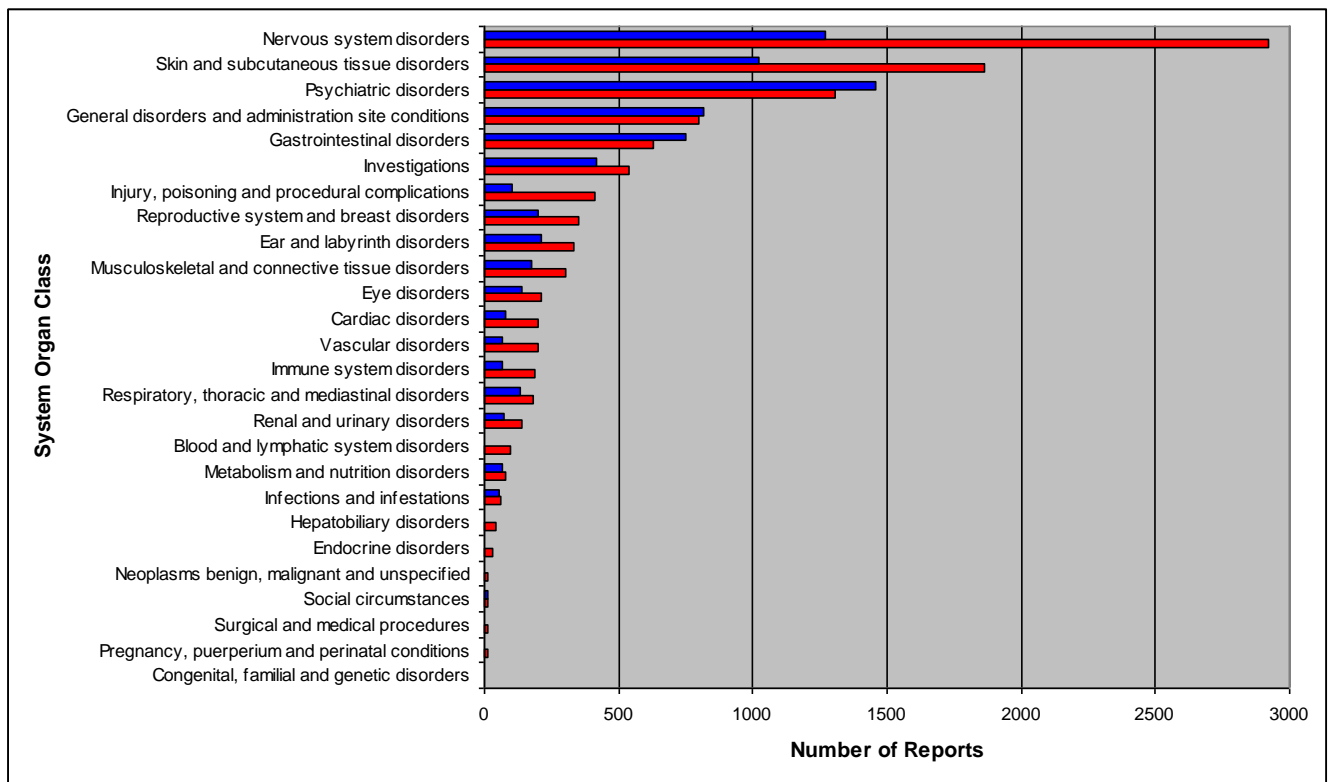


Figure 9a – The upper bar of each pair represents numbers of reports from Consumers (blue) and the lower bar reports from Health Care Professionals (red) (Population 1)



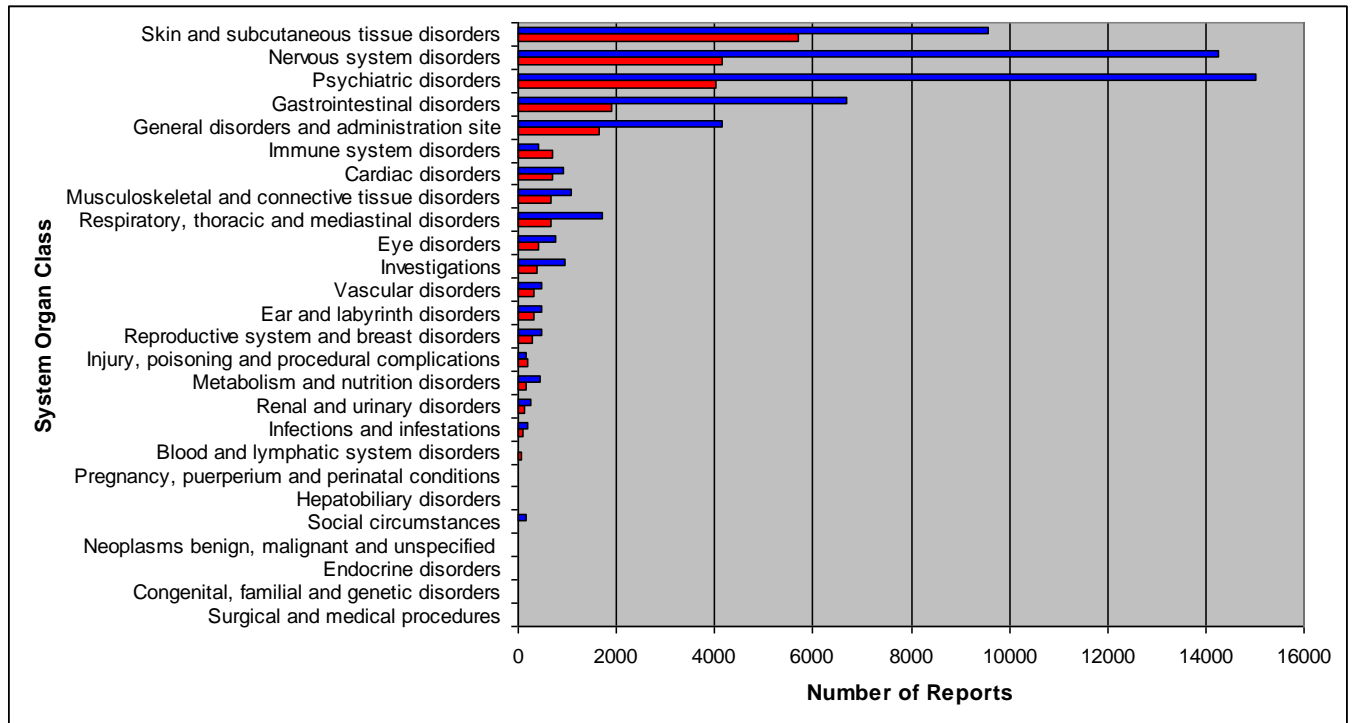


Figure9b – The upper bar of each pair represents numbers of reports from Consumers (blue) and the lower bar reports from Health Care Professionals (red) (Population 2)

Most Frequent On-Therapy Adverse Events  
 PTs sorted by relative risk

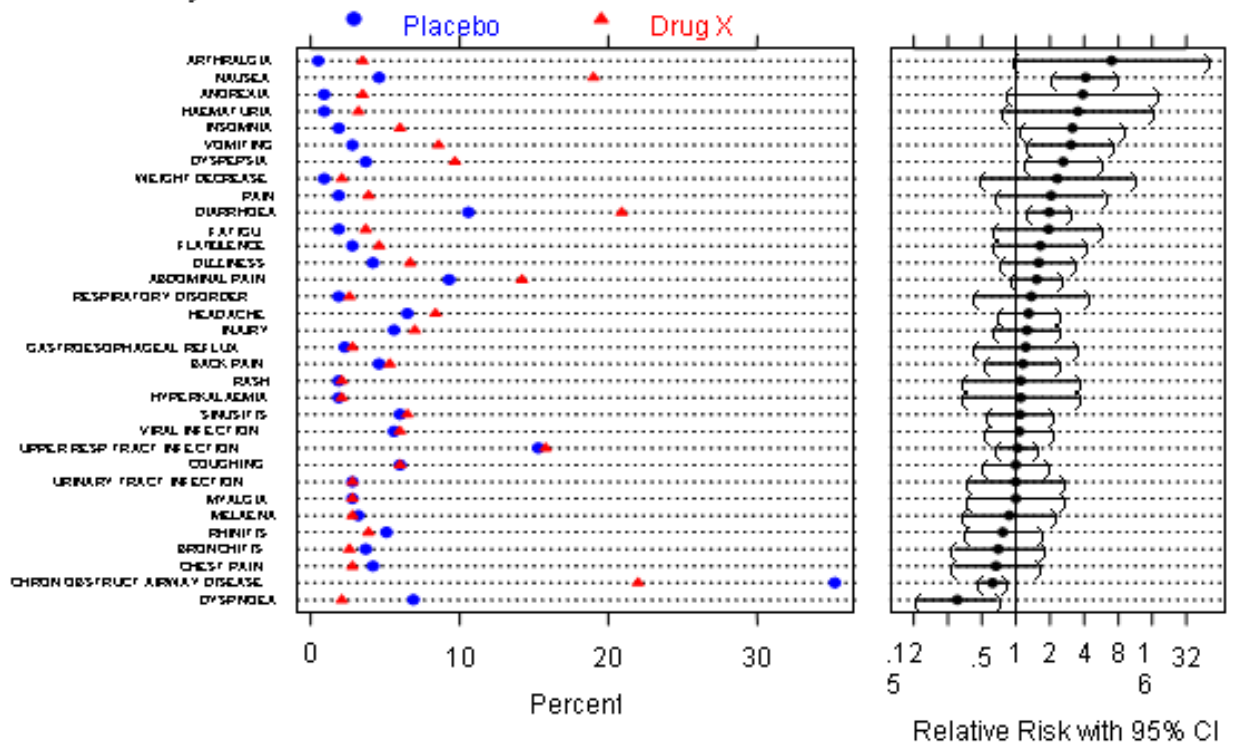


Figure 10 – For a small dataset, a display of PTs may be adequate

## SOC Infections and infestations

### Primary SOC Analysis

Adverse Event (MedDRA v17.0)	25 mg MyDrug (N=44)	Placebo (N=15)
SOC Infections and infestations	14 (31.8%)	4 (26.7%)
PT Upper respiratory tract infection	5	2
PT Sinusitis	3	0
PT Urinary tract infection	2	1
PT Ear infection	2	0
PT Viral infection	2	0
PT Bronchitis	1	0
PT Influenza	1	0
PT Localised infection	0	1
PT Lower respiratory tract infection	1	0
PT Pneumonia	1	0
PT Tooth abscess	1	0

### Secondary SOC Analysis (same data as above)

Adverse Event (MedDRA v17.0)	25 mg MyDrug (N=44)	Placebo (N=15)
SOC Respiratory, thoracic and mediastinal disorders		
PT Upper respiratory tract infection	5	2
PT Sinusitis	3	0
PT Bronchitis	1	0
PT Influenza	1	0
PT Lower respiratory tract infection	1	0
PT Pneumonia	1	0
SOC Infections and infestations		
PT Viral infection	2	0
PT Localised infection	0	1
SOC Renal and urinary disorders		
PT Urinary tract infection	2	1
SOC Ear and labyrinth disorders		
PT Ear infection	2	0
SOC Gastrointestinal disorders		
PT Tooth abscess	1	0

Figure 11 – Programmed primary and secondary SOC outputs

Asthma/bronchospasm (SMQ) Cases – Narrow Search  
(since 1-JAN-2008)

ID	MedDRA_PT	REPORT_VERBATIM	DATE_CREATED
045	Asthma	Asthma attack	01-APR-2008
063	Asthma	Severe asthma	10-JUN-2008
060	Asthma exercise induced	Asthma when exercising	30-MAY-2008
091	Bronchospasm	Spasms, bronchial	12-AUG-2008
074	Bronchospasm	Bronchoconstriction	03-JUL-2008
100	Bronchial hyperreactivity	Airways hyperreactive	20-SEP-2008
069	Bronchial hyperreactivity	Reactive airways disease	21-JUN-2008

Asthma/bronchospasm (SMQ) Cases – Broad Search  
(since 1-JAN-2008)

ID	MedDRA_PT	REPORT_VERBATIM	DATE_CREATED
023	Allergic respiratory disease	Respiratory (allergy) disorder	18-FEB-2008
045	Asthma	Asthma attack	01-APR-2008
063	Asthma	Severe asthma	10-JUN-2008
060	Asthma exercise induced	Asthma when exercising	30-MAY-2008
016	Bronchial obstruction	Bronchial obstruct.	16-JAN-2008
039	Bronchial obstruction	Bronchus obstruction	14-MAR-2008
091	Bronchospasm	Spasms, bronchial	12-AUG-2008
074	Bronchospasm	Bronchoconstriction	03-JUL-2008
100	Bronchial hyperreactivity	Airways hyperreactive	20-SEP-2008
069	Bronchial hyperreactivity	Reactive airways disease	21-JUN-2008
088	Obstructive airways disorder	Obstructive airways disorder	29-JUL-2008
049	Obstructive airways disorder	Obstructed airways dis.	20-APR-2008
022	Wheezing	Wheeze	16-FEB-2008
031	Wheezing	Wheezes	02-MAR-2008
106	Wheezing	Wheezing	28-SEP-2008
046	Wheezing	Wheezing (acute)	06-APR-2008

*Figure 12 – Results of Narrow and Broad SMQ Searches*