



Shared Health Summary My Health Record Usability Recommendations

16 June 2017 V1.3 Approved for external use Document ID: DH-2211:2017 Australian Digital Health Agency

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Acknowledgements

Council of Australian Governments

The Australian Digital Health Agency is jointly funded by the Australian Government and all state and territory governments.

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Document information

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Product version history

Product version	Date	Release comments
1.0	25 Nov 2013	Extracted from Clinical Usability Program (CUP) R1 PCEHR Clinical Usability Software Requirements v1.0
1.1	5 May 2014	Revised version, incorporating usability recommendations from both CUP R1 and CUP R2
1.2	31 Dec 2014	Revised version, incorporating usability recommendations from CUP R3 and minor editorial updates.
1.3	16 Jun 2017	Revised version, incorporating minor amendments to existing recommendations to align with the latest <i>Shared Health Summary</i> <i>Structured Content Specification</i> . Editorial updates related to renaming of PCEHR system to My Health Record system. Rebrand as Agency document.

Table of contents

1	Introd	luction	5	
	1.1	Purpose	5	
	1.2	Scope	5	
	1.3	Conformance	5	
	1.4	How to use this document	6	
2	Attesting a shared health summary			
3	Popula	ating a shared health summary1	0	
4	Prese	ntation and selection of medical history items1	13	
5	Medic	al history narrative 1	14	
6	Exclus	sion statements for medical history items1	15	
Glos	sary	1	8	
Defe	ossary			

1 Introduction

1.1 Purpose

It has been recognised that developers of software systems that access the My Health Record system¹ need usability recommendations to complement the software requirements provided by other eHealth specifications. These usability recommendations are designed to achieve greater consistency between general practice software products that access the My Health Record system, thereby improving clinical usability.

While the usability recommendations are developed specifically for general practice software vendors, they are provided to all software developers interested in improving the usability of their software systems. They are *not* part of the set of software conformance requirements for clinical information systems accessing the My Health Record system; however, conformance to these usability recommendations is strongly encouraged by clinical system users. More information about conformance to the usability recommendations is provided in section 1.3.

These recommendations were prepared as part of Agency's Clinical Usability Program (CUP) in consultation with general practice clinicians.

As the My Health Record system functionality increases, these usability recommendations will be updated.

1.2 Scope

This document provides usability recommendations for clinical information systems and contracted service provider systems authoring or rendering information contained in shared health summary documents exchanged with the My Health Record system.

It is focused on recommendations applicable specifically to shared health summary documents. Additional usability recommendations for *all* types of clinical documents are published in the *Clinical Documents - My Health Record Usability Recommendations v1.3.*²

This document does *not* provide usability recommendations for:

- document types other than shared health summary;
- My Health Record system functions not related to the authoring and rendering of shared health summary documents exchanged with the My Health Record system; or
- display and management of clinical terminology.

1.3 Conformance

Software developers may want to claim that their software implements these usability recommendations. For such claims to be meaningful and for the healthcare community to have a shared understanding of these claims, the usability recommendations have been documented in the form of software conformance

¹ Previously known as the personally controlled electronic health record (PCEHR) system.

² Links to this and other referenced documents are given in the References section at the end.

requirements using the standard conformance verbs **SHALL**, **SHALL NOT**, **SHOULD** and **SHOULD NOT**.

Conformance to the recommendations in this document is not a prerequisite for software to be granted access to the My Health Record system. However, conformance to these usability recommendations is strongly encouraged by clinical software users. A software developer wanting to claim conformance to the clinical usability recommendations must have software that conforms to all mandatory and applicable conditional recommendations in this document. These are the recommendations using the verbs **SHALL** and **SHALL NOT**. Achievement of conformance to the mandatory and applicable conditional recommendations is recognised by mention on the table titled "eHealth Functions Available – General Practice Software Products" on the Agency website.

It is expected that the vendor's software will still meet (or has met) the requirements listed in the *Shared Health Summary PCEHR Conformance Profile* v1.6.

1.4 How to use this document

How you use this document depends on whether you have used either of the previous versions of this document, which incorporated CUP releases 1, 2 and 3:

- Version 1.1 incorporating CUP releases 1 and 2 (R1 and R2). CUP R1 recommendations focused on usability in the creation of a shared health summary document and the display of the My Health Record document list, while CUP R2 focused on the creation of an event summary document. Version 1.1 of this document should be read in conjunction with *Clinical Documents PCEHR Usability Recommendations v1.1* and *Event Summary PCEHR Usability Recommendations v1.0*
- Version 1.2 incorporating CUP release 3 (R3). These recommendations focused on the creation of a 'home page' when the user opens a patient's digital health record and also on identifying and indicating new clinical documents in a patient's digital health record to the user. Version 1.2 of this document should be read in conjunction with *Clinical Documents PCEHR Usability Recommendations v1.2* and *Event Summary PCEHR Usability Recommendations v1.1*

Vendors who have *not* implemented any CUP recommendations are advised to implement all recommendations listed in this document, to increase usability of their software.

Vendors who have implemented previous releases of CUP recommendations are advised to review the 'status' column of each recommendation to determine whether they will need to implement or not, based on the following guidelines:

- No change from v1.1 this status means that the recommendation was introduced in either R1 or R2 with no further amendments
- No change from v1.2 this status means that the recommendation was introduced in R3 with no further amendments
- **Revised for v1.2** this status means that the recommendation was amended in v1.2 but remains unchanged in v1.3
- **Revised for v1.3** this status means the recommendation has been amended in v1.3

For details of the changes made to recommendations between the various versions, refer to the *CUP Usability Recommendations – Complete Change Log v1.0*, available on request from the Agency Help Centre.

The shared health summary specifications have been updated to incorporate a number of CUP recommendations – these are noted throughout this document. If your software conforms to the updated specifications, then those CUP recommendations that have been incorporated are not applicable.

Some recommendations have had editorial changes, for example to change *PCEHR* to *digital health record*, and *PCEHR system* to *My Health Record system*. These changes do not alter the meaning of the recommendation, and are marked as 'no change'.

2 Attesting a shared health summary

Applies to: General practice document authoring systems (shared health summary clinical documents).

The *Shared Health Summary PCEHR Conformance Profile* (conformance profile) states that:

A clinical information system shall display the final version of a shared health summary to the author and prompt the author to attest to the content of the shared health summary before the clinical information system uploads the shared health summary to the My Health Record system and to assert the healthcare provider individual (i.e. the author of the shared health summary) is a nominated healthcare provider as defined by the *Personally Controlled Electronic Health Records Act 2012.*³ [COM2012]

There are multiple ways to implement this requirement. The conformance profile states:

One option for meeting this requirement is for a clinical information system to display the shared health summary along with a user interface button with the statement "By uploading this Shared Health Summary, I acknowledge that I am a Nominated Healthcare Provider for this patient as defined by the *Personally Controlled Electronic Health Records Act.*

Software vendors have taken a variety of approaches to implementing these requirements; some have introduced extra steps or checks concerning accuracy and patient consent.

There is no legal requirement for a provider to obtain the consent of a consumer each time a shared health summary is uploaded, as the provider can rely on the standing consent given by the consumer at registration.

The following professionals are eligible to be nominated healthcare providers as defined by the *My Health Records Act*:

- medical practitioners
- registered nurses
- Aboriginal and/or Torres Strait Islander health practitioners (with a Certificate IV in Aboriginal and/or Torres Strait Islander Primary Health Care (Practice)

The shared health summary authoring software is not required to enforce that the author falls into one of these categories, and doing so reliably may not be possible.

The usability recommendations in this section address how the shared health summary authoring review should be presented to the user.

³ This has now been renamed to My Health Records Act 2012 (Cth). Both names appear in this document.

ID	Recommendation	Obligation	Status
SHS.01	The software SHALL present a statement on shared health summary review and attestation as follows:	Mandatory	No change from v1.1
	 I am the patient's nominated healthcare provider in accordance with the <i>My Health Records Act 2012</i>. I am providing ongoing care to this patient. I have prepared this shared health summary in consultation with the patient. 		
	Note: The above wording meets the requirements of the conformance process, addresses and encourages best clinical practice as agreed by clinicians consulted during the Clinical Usability Program, and does not intrude on the natural workflow of a clinician.		
	The wording does not ask the clinician to confirm that the document is either complete or accurate, since the patient may ask to withhold information from the document.		
SHS.02	The software SHALL have one and only one confirmation step for shared health summary review and attestation.	Mandatory	No change from v1.1
	Note: The recommendation reduces the key strokes needed to submit a shared health summary. A single upload button is sufficient to meet this recommendation. It is not necessary to display a checkbox to record consent, nor is it necessary to display the acknowledgement as a separate pop-up window.		
SHS.03	The software SHALL NOT upload the authored clinical document without an explicit confirmation from the user.	Mandatory	No change from v1.1
	Note: A generic keystroke such as "Tab" or "Enter" is liable to be pressed automatically by the user, so it would not count as an explicit confirmation. Examples of explicit confirmations include a mouse click on an "Upload" button, or a dialog with "Y" or "N" responses.		

3 Populating a shared health summary

Applies to: General practice document authoring systems (shared health summary clinical documents).

When creating a shared health summary, it should be possible for the clinician to readily manage the relevant clinical data items to be included. Current implementations typically present lists of data items drawn from the patient record with a check box against each list item denoting inclusion or exclusion in the shared health summaries. Each list corresponds to a health summary document section – that is, adverse reactions, medications, medical history or immunisations.

The methods and rules that determine the initial check box state (that is, whether a data item is included by default) have not been specified. Consequently, there are considerable differences between software products. Various inconsistent approaches based on clinical entry "shared data flags" (for example, include in summary, include in correspondence) and direct selection on authoring have been implemented in systems.

The *Shared Health Summary Structured Content Specification v1.2* provides information about the sections of a shared health summary. In brief:

- Adverse Reactions: Information about adverse reactions or propensity to adverse reaction of the patient (including allergies and intolerances), and any relevant reaction details.
- *Medications:* Medicines which the healthcare consumer is using. This includes self-prescribed, clinician-prescribed and non-prescription medicines. This section must not be used to record vaccine administration records of the healthcare consumer. The "Administered immunisation" section must be used for this purpose.
- *Medical History:* The past and current medical history of the healthcare consumer which is relevant to the episode of care. This includes problem/diagnosis and medical or surgical procedures performed.
- *Immunisations:* Information about the immunisation history of the healthcare consumer.

ID	Recommendation		Status	
SHS.04	The software SHALL display four complete lists of data items sourced from the local patient record:	Mandatory	No change from v1.1	
	1 Adverse Reactions			
	2 Medications			
	3 Medical History			
	4 Immunisations			
	The list headings SHALL be as given above.			
	Information in these lists SHALL be in accordance with the definition of Adverse Reactions, Medications, Medical History and Immunisations, stated in in the <i>Shared Health Summary Structured Content Specification</i> .			

ID	Recommendation	Obligation	Status
SHS.05	If the software allows an individual data item to be attributed with a "Confidentiality" flag and the flag is set to "true", then the software SHALL disallow the selection and inclusion of that data item.	Conditional	No change from v1.1
	Note: Some clinical systems incorporate confidentiality settings that restrict sharing of sensitive information or particular health information (for example, HIV status) outside the healthcare organisation. Such items will still appear in the shared health summary review authoring screen; however, their status should be clearly indicated to the user. Furthermore, the software may implement controls to prevent inadvertent inclusion in a shared health summary by disabling selection when authoring a shared health summary.		
SHS.06	If the software has one or more item-level shared data flags for an entry type, those items that have been flagged for sharing SHALL be marked for inclusion in the shared health summary by default.	Conditional	No change from v1.1
	Note: Entry types are: adverse reactions, medications, medical history and immunisations. Clinical systems commonly attribute medical history entries with shared data flag(s). When authoring a shared health summary, it must be possible to de-select any item marked for default inclusion.		
SHS.07	If the software does not have an item-level shared data flag for an entry type, all items of that entry type SHALL be marked for inclusion by default.	Conditional	No change from v1.1
	Note: Clinical systems typically do not attribute adverse reactions, medications and immunisations with shared data flags. When authoring a shared health summary, it must be possible to de-select any item marked for default inclusion.		
SHS.08	The software SHALL support selection and de-selection of clinical items for shared health summaries, regardless of whether the shared data flag is marked or unmarked on entry.	Mandatory	No change from v1.1

ID	Recommendation	Obligation	Status
SHS.21	When authoring a new shared health summary, the software SHALL pre-select items that were previously included in the most recently authored shared health summary for the patient uploaded by the user's healthcare organisation, unless the item has since been removed or has been marked as inactive or confidential.	Mandatory	No change from v1.2
	Note: In addition to this recommendation, any new items will be pre-selected as set out in recommendations SHS.05, SHS.06 and SHS.07. Recommendation SHS.08 allows the user to override any pre-selections before uploading the shared health summary.		
	This recommendation allows users to avoid having to reselect items to add to a new shared health summary if the item was previously included in the last shared health summary uploaded by the same organisation and the item has not since been removed or marked as inactive or confidential.		
	Some vendors may have taken the initiative and implemented this recommendation already in their software.		

4 Presentation and selection of medical history items

Applies to: General practice document authoring systems (shared health summary clinical documents).

The clinical requirements for the Medical History Section (as specified by the *Shared Health Summary Information Requirements* v1.1) describe a single list of medical history items, with a single exclusion statement. However the *Shared Health Summary CDA Implementation Guide* v1.4 separates medical history into three sub-groupings: Problems/Diagnosis, Procedures, and Medical History Items. There are two exclusion statements: one for Problems/Diagnosis and another for Procedures. The need for two statements is due to the way the information is modelled and handled internally – it was not intended to impact on the way that the information guide does not directly address the question of how documents are presented and, since it suggests separate lists, this is how the shared health summary has been implemented.

Information on a patient's medical history is to be presented to the user so that they can select the items to record in a shared health summary.

ID	Recommendation	Obligation	Status
SHS.09	The software SHALL present to the user a combined medical history list containing all applicable procedures, problems/diagnosis and uncategorised medical history entries.	Mandatory	Revised for v1.3
SHS.10			Revised for v1.3
	Note: It is not expected that the software presents history items contained in progress notes or similar narrative entries.		

5 Medical history narrative

Applies to: General practice document authoring systems (shared health summary clinical documents).

The Shared Health Summary CDA Implementation Guide states that the shared health summary medical history section has one narrative for problems/diagnosis, procedures and medical history items and three sub-sections for the structured data for problems/diagnosis, procedures and medical history items. The intent is for the medical history narrative to be formatted as a table according to the recommendations listed below.

ID	Recommendation	Obligation	Status	
SHS.11	The software SHALL populate the narrative of the medical history section of the CDA document with a single table containing all procedures, problem/diagnoses or other medical histories that are to be included in the document.	Mandatory	No change from v1.1	
SHS.12	The software SHALL create a medical history table with an "Item" column containing a textual description of the problem/diagnosis, procedure or uncategorised medical history item.	Mandatory	Revised for v1.3	
SHS.13	If the software supports a date entry associated with medical history items, then the software SHALL create a medical history table with a "Date" column containing a point in time or time period for the entry.	Conditional	No change from v1.1	
	Note: The date displayed could be:			
	 a specific date corresponding to a point in time occurrence, such as a procedure date or date of onset; 			
	 a date range, such as date of onset to date of remission; or 			
	 "(ongoing)" to indicate that the condition is ongoing. 			
SHS.14	If comments associated with medical history items are supported, then the software SHALL create a medical history table with a "Comment" column containing additional comments about the entry.	Conditional	No change from v1.1	
SHS.15	The software SHALL create entries in the medical history table in reverse chronological order based on the Date field. Medical history items with the most recent dates SHALL be listed higher in the table.	Mandatory	No change from v1.1	
	Any medical history items with no corresponding date (e.g. where no date of onset has been recorded) SHALL be displayed at the top of the table.			
	Note: When a history item has a date range recorded (e.g. date of onset to date of resolution), the start of the date period is used for sorting.			

6 Exclusion statements for medical history items

Applies to: All general practice systems authoring shared health summaries that record exclusion statements.

The medical history section of shared health summary clinical documents allows three possible types of medical history entry (problem/diagnosis, procedure and uncategorised medical history item) and two types of global exclusion statements (for problems and diagnoses and for procedures) as shown below.

Medical History Entry

Global Exclusion Statement

- Problem/Diagnosis
- Exclusion Statement Problems and Diagnoses

Procedure

- Exclusion Statement Procedures
- Uncategorised Medical History Item
 Exclusion Statement -

A clarified definition of "Uncategorised Medical History Item" based on the *Shared Health Summary Structured Content Specification* definition is:

"A medical history entry which cannot be categorised into one of the categories such as Procedure and Problem/Diagnosis."

This covers cases where the source system cannot automatically classify an entry as a Problem/Diagnosis or a Procedure, including cases where:

- The coding system used for medical history item cannot structurally support adequate concept classification.
- The medical history item is maintained as free-text and thus has never been classified.

Appropriate use is defined here based on the existence of medical history entries of each type.

Since it is not known whether an "Uncategorised Medical History Item" entry is conceptually a procedure or a problem/diagnosis, exclusion statements cannot be used when an "Uncategorised Medical History Item" entry is present, as the entry may, in fact, be a procedure or a problem/diagnosis. Therefore, the following explicit constraints apply based on the *Shared Health Summary CDA Implementation Guide*.

	Recommendation				Obligation	Status		
SHS.16	When there are no entries for both "Procedure" and "Uncategorised Medical History Item" the software SHALL create an "Exclusion Statement – Procedures".				Mandatory	Revised for v1.3		
	Example:							
	Medical History Entry		Global Exclusion Stater	nent				
	Problem/Diagnosis		Problems and Diagnoses					
	Procedure	No entry	Procedures	Required				
	Other Medical History It	em No entry						
	<i>implemented aga</i> Structured Conte	<i>ninst either i</i> nt Specifica	<i>is not applicable if t</i> <i>the</i> Shared Health S tion v1.2 <i>or later O</i> lementation Guide v	Summary – <i>R the</i> Shared				
SHS.17	"Uncategorised N	ledical Histo	r both "Problem/Dia ory Item" the softwa ent – Problems and	are SHALL	Mandatory	Revised for v1.3		
	Example:							
	Medical History Entry		Global Exclusion State	ment				
	Problem/Diagnosis	No entry	Problems and Diagnoses	Required				
	Procedure		Procedures					
	Other Medical History It	em No entry						
				Note: This recommendation is not applicable if the system has implemented against either the Shared Health Summary – Structured Content Specification v1.2 or later OR the Shared Health Summary – CDA Implementation Guide v1.4 or later.				
	<i>implemented aga</i> Structured Conte	<i>ninst either i</i> nt Specifica	the Shared Health S tion v1.2 or later O	Summary – R the Shared				
5HS.18	implemented aga Structured Conte Health Summary When there are n Diagnosis" and "U software SHALL	ninst either a nt Specifica – CDA Imp no entries fo Jncategorise create both	the Shared Health S tion v1.2 or later O	Summary – <i>R the</i> Shared v1.4 <i>or later.</i> , "Problem/ tem" the sement –	Mandatory	Revised for v1.3		
SHS.18	implemented aga Structured Conte Health Summary When there are r Diagnosis" and "U software SHALL Procedures" and Diagnoses".	nt Specifica – CDA Imp o entries fo Jncategorise create both "Exclusion S	the Shared Health S tion v1.2 or later O lementation Guide r all of "Procedure" ed Medical History I an "Exclusion State	Summary – <i>R the</i> Shared v1.4 <i>or later.</i> , "Problem/ tem" the ement – ms and	Mandatory			
5HS.18	implemented aga Structured Conte Health Summary When there are r Diagnosis" and "U software SHALL Procedures" and Diagnoses". Example:	nt Specifica – CDA Imp o entries fo Jncategorise create both "Exclusion S	the Shared Health S tion v1.2 or later O lementation Guide v r all of "Procedure" ed Medical History I an "Exclusion State Statement – Probler	Summary – <i>R the</i> Shared v1.4 <i>or later.</i> , "Problem/ tem" the ement – ms and	Mandatory			
SHS.18	implemented aga Structured Conte Health Summary When there are r Diagnosis" and "U software SHALL Procedures" and Diagnoses". Example: Medical History Entry	nt Specifica – CDA Imp o entries fo Jncategorise create both "Exclusion S	the Shared Health S tion v1.2 or later O lementation Guide v r all of "Procedure" ed Medical History I an "Exclusion State Statement – Probler	Summary – <i>R the</i> Shared v1.4 <i>or later.</i> , "Problem/ tem" the ement – ms and ment	Mandatory			

ID	Recommendation				Obligation	Status
SHS.19	When there are any entries for "Uncategorised Medical History Item" the software SHALL NOT create an exclusion statement.			Mandatory	Revised for v1.3	
	Example:					
	Medical History Entry	Global Exclusion Statem	ent	-		
	Problem/Diagnosis	Problems and Diagnoses	Not allowed	-		
	Procedure	Procedures	Not allowed	-		
	Other Medical History Item Entry			-		
	be an exclusion statement Note: This recommendation implemented against either Structured Content Specific Health Summary – CDA Im					
SHS.20	The software SHALL NOT use the "not asked" exclusion statement in the context of Shared Health Summary clinical document authoring.				Mandatory	No change from v1.1
	Note: This recommendation implemented against either Structured Content Specific Health Summary – CDA Im	<i>the</i> Shared Health S ation v1.2 <i>or later O</i>	Summary – <i>R the</i> Share	d		

Note: In the example tables above, the blank cells in Medical History Entry indicate either an entry or blank. A medical history section is allowed to contain procedures, problem/diagnosis, and other uncategorised medical history items.⁴ Having both categorised items (procedures and problem/diagnosis) and uncategorised medical history items would be unusual, because generally if a system is able to differentiate some items, it is able to differentiate them all. However a system may be able to categorise some, and not others – because of legacy data, or partial classification in the underlying terminology, for instance. For this reason, the rules allow a mix of categorised and uncategorised items.

⁴ In the section "Clarifications" of the *Shared Health Summary Release Note v1.5*, the document states to "use EITHER "Problem/Diagnosis" and "Procedure" OR "Uncategorised Medical History Item", but NOT both". This should be understood as product guidance and is consistent with these usability recommendations.

Glossary

Term	Description
Clinical Document Architecture (CDA)	An XML-based mark-up standard intended to specify the encoding, structure and semantics of clinical documents exchanged between health software systems. Specifications for clinical documents are based on CDA Release 2 [HL72005].
clinical document	A digital file containing personal health information about an individual, containing unstructured (narrative) information and optionally structured (atomic) information.
clinical information system (CIS)	A system that deals with the collection, storage, retrieval, communication, and use of health related data, information and knowledge pertaining to subjects of care [AS5021]. The system may comprise one or more applications or components.
contracted service provider	An entity that may offer health software as a service, and support access to the My Health Record system on behalf of healthcare organisations. A contracted service provider provides under a contract with the healthcare provider organisation: a) information technology services relating to the My Health Record system; or b) health information management services relating to the My Health Record system (Section 5 <i>My Health Records Act 2012</i>).
healthcare consumer	A person who is the subject of care.
My Health Record system	National eHealth infrastructure for managing records in eHealth. This was previously called the personally controlled electronic health record system (eHealth record system).

References

Australian Digital Health Agency references

The references below are published on <u>www.digitalhealth.gov.au</u>.

If viewing this as a printed document, use the DH-XXXX:YYYY for new documents and NEHTA-XXXX:YYYY for old documents as identifier to search for the reference online.

<u>NEHTA-1840:2015</u>	Shared Health Summary CDA Implementation Guide v1.4, 10 April 2015
<u>NEHTA-1837:2015</u>	Shared Health Summary Information Requirements v1.1, 10 April 2015
<u>NEHTA-1839:2015</u>	Shared Health Summary Structured Content Specification v1.2, 10 April 2015
<u>NEHTA-1852:2015</u>	Shared Health Summary Release Note v1.5, 10 April 2015
NEHTA-1838: 2015	Shared Health Summary PCEHR Conformance Profile v1.6, 10 April 2015
<u>NEHTA-2210:2016</u>	<i>Clinical Documents My Health Record Usability Recommendations v1.3,</i> 6 January 2016

Other references

At the time of publication, the versions below were valid. However, as all documents are subject to revision, readers are encouraged to use the most recent versions of these documents.

[AS5021]	AS 5021:2005 - The language of health concept representation, Standards Australia, 2005.
[COM2012]	My Health Records Act 2012 (previously named the Personally Controlled Electronic Health Records Act 2012), Australian Government ComLaw, 2012
[HL72005]	Clinical Document Architecture, Release 2, ISO/HL7 27932:2008, 21 Apr 2005