

eHealth Pathology Report Release Note

28 February 2021 v1.2.3 Approved for external use Document ID: DH-3403:2021

Related end product identifier: EP-3399:2021

Release rationale

This incremental release provides a replacement update for the template packages of the eHealth Pathology Report document type.

Previous versions of the template packages performed extensive validations of field values against hard-coded terminologies. These terminology validations have now been removed where:

- Terminologies change over time (template packages only perform static validations)
- Terminologies provide limited value

This allows for the successful upload of eHealth Pathology Report documents where the value of having the clinical information in the My Health Record system clearly outweighs any minor terminology misalignments.

Please refer to the Change Details section below for further details.

Package inclusions

New

None

Updated (supersedes previous version)

Identifier	Name and version	
DH-3403:2021	eHealth Pathology Report – Release Note v1.2.3 (this document)	
DH-3395:2021	eHealth Pathology Report – Template Package Library v1.1.4	

No change

Identifier	Name and version	
DH-2525:2017	eHealth Pathology Report – Conformance Test Specification v1.0	
NEHTA-1412:2014	Pathology Report – CDA $^{\mathrm{m}1}$ Implementation Guide v1.0	
NEHTA-1413:2014	Pathology Report — Structured Content Specification v1.0	
NEHTA-1884:2014	eHealth Pathology Report – Information Requirements v1.1	
NEHTA-2214:2016	eHealth Pathology Report – My Health Record Conformance Profile v1.1	

Removed (archived or withdrawn)

None

¹ CDA is a trademark of Health Level Seven International and is registered with the United States Patent and Trademark Office.

Change details

Validations for the following terminologies have been removed from the template packages of this document type:

- TelecommunicationAddressUse
- Health Care Client Name Usage
- Health_Care_Provider_Organisation_Name_Usage
- AS_5017-2006_Health_Care_Client_Source_of_Death_Notification
- Health_Care_Client_Identifier_Address_Purpose
- Health Care Client Electronic Communication Medium
- CDA-Act.moodCode
- HL7_ObservationInterpretationNormality
- Australian_Postal_Delivery_Number
- Australian_Postal_Delivery_Type_Code
- Australian_Street_Suffix_Code
- Australian_Unit_Type
- ISO3166-1 (Countries)
- 0 ANZSCO Australian and New Zealand Standard Classification of Occupations, First Edition, Revision 1
- 0 ANZSCO Australian and New Zealand Standard Classification of Occupations, First Edition,
 2006
- Coded_DataType_NullFlavor

The template package library now contains the following template packages:

Document type variant	Conformance level	Template package ID
HPIIRelaxed	3A	1.2.36.1.2001.1006.1.220.3 - CL 3A (svn-40639)
Default	3A	1.2.36.1.2001.1006.1.220.4 - CL 3A (svn-40639)

Systems uploading a clinical document to the My Health Record system need to provide the correct template package identifier. The full list of published template packages can be found in the *Template Package Directory* v1.14².

Known issues

None

² https://developer.digitalhealth.gov.au/resources/clinical-documents-template-package-directory-v1-14

Stakeholders

The following stakeholders were engaged for the development of this release:

- Developers of systems generating eHealth Pathology Report documents;
- My Health Record System Operator;
- National Infrastructure Operator.

Audience

The intended audience of this document includes:

- Developers of systems generating eHealth Pathology Report documents;
- My Health Record System Operator;
- National Infrastructure Operator.
- Senior managers and policy makers, support teams and system integrators.

Support

For further support or to provide feedback, please email help@digitalhealth.gov.au

Previous releases

Date Version

21 Dec 2017

EP-2558:2017 eHealth Pathology Report v1.2.2

Release Note

Release rationale

This incremental release provides a number of defect fixes for the template packages of the eHealth Pathology Report document type. For the majority of developers, there is no need to update their systems, as the updated template packages have the same template package identifiers as the previous ones.

However, developers should review whether their systems are affected by one defect fix that is not backwards compatible. The highest level "REPORTING PATHOLOGIST" element of eHealth Pathology Report documents is now validated as mandatory. This change brings the template packages in line with the published specifications but may require updates to systems that currently don't provide the "REPORTING PATHOLOGIST" information.

16 Jun 2017

EP-2454:2017 eHealth Pathology Report v1.2.1

Release note

Release rationale

This release of the eHealth Pathology Report end product provides updates to the eHealth Pathology Report Template Package Library and new eHealth Pathology Report Conformance Test Specification.

eHealth Pathology Report Template Package Library v1.1.2

The Template Package Library includes defect fixes for template packages without HPI-I relaxation. The default (i.e. non-relaxed) template packages for eHealth Pathology Report document types has been updated to allow the omission of HPI-Is for the document author and the reporting pathologist in case of the document having been authored by more than one pathologist.

The template package library contains the following template packages:

Document type variant	Conformance level	Template package ID
HPIIRelaxed	3A	1.2.36.1.2001.1006.1.220.3 - CL 3A (svn-37049)
Default	3A	1.2.36.1.2001.1006.1.220.4 - CL 3A (svn-39116)

Document producers need to ensure they provide corresponding template package IDs when uploading documents to the My Health Record system. The full list of published template packages can be found in the <u>Clinical Documents - Template Package Directory v1.8</u>.

 $e Health\ Pathology\ Report\ Conformance\ Test\ Specification\ v1.0$

This is an initial release of the eHealth Pathology Report Conformance Test Specification. This Conformance Test Specification consists of a series of test cases, each of which is a discrete test and a statement of expected behaviour. Vendors can use the documented test cases to assess the conformance of their Pathology eHealth Software.

10 Mar 2016

EP-2242:2016 eHealth Pathology Report v1.2

Release note

Release rationale

This release of the eHealth Pathology Report end product updates the following product components:

- My Health Record conformance profile
- Template package library.

The consent management section of the conformance profile (Section 3.4) has been rewritten to better reflect implementation requirements as a result of feedback from early adopters. Section 3.5 has updates to record keeping requirements to allow greater flexibility is how records are kept.

Document producers need to ensure they provide corresponding template package IDs when uploading documents to the My Health Record system. The full list of published template packages can be found in the <u>Clinical Documents - Template Package Directory v1.6</u>.

31 Jul 2015

EP-2050:2015 eHealth Pathology Report v1.1

Release note

Release rationale

This release of the eHealth Pathology Report end product adds the personally controlled electronic health record (PCEHR³) conformance profile and associated updates to the template package library.

The eHealth Pathology Report PCEHR Conformance Profile summarises the requirements for producers and consumers of eHealth Pathology Reports that connect to the PCEHR system.

The template package library contains the following template packages. Document producers need to ensure they provide corresponding template package IDs when uploading documents to the PCEHR system.

Document type variant	Conformance level	Template package ID
HPIIRelaxed	3A	1.2.36.1.2001.1006.1.220.3 - CL 3A (svn-37049)
Default	3A	1.2.36.1.2001.1006.1.220.4 - CL 3A (svn-37049)

The full list of published template packages can be found in the *Template Package Directory v1.5*. This end product has a dependency on *Clinical Documents - Common Conformance Profile v1.6*.

31 Dec 2014

EP-1882:2014 eHealth Pathology Report v1.0

Release note

Release rationale

This is the initial release of the eHealth Pathology Report specifications.

The eHealth Pathology Report specification set (end product) provides an enhancement to the PCEHR system. It includes a CDA document containing a pathology report that can be uploaded to the PCEHR system. Individuals and healthcare providers with access to a person's PCEHR will be able to retrieve Pathology Report CDA documents through the individuals PCEHR using the PCEHR portals or local clinical information systems via the B2B Gateway.

This set of specifications is accompanied by an eHealth Pathology Report View (published as a separate end product EP-1982:2014) which allows listing, grouping and sorting of pathology reports in the individual's PCEHR.

The eHealth Pathology Report specifications form part of the foundational set of specifications to support the development of an individual's PCEHR.

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³ Clarification: PCEHR means the My Health Record, formerly the "Personally Controlled Electronic Health Record", within the meaning of the *My Health Records Act 2012* (Cth), formerly called the *Personally Controlled Electronic Health Records Act 2012* (Cth).

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