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# **Dispense Record - Structured Document Template**

## **Release Note**

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Release: 1

Public

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**National E-Health Transition Authority Ltd**

Level 25

56 Pitt Street

Sydney, NSW, 2000

Australia.

[www.nehta.gov.au](http://www.nehta.gov.au)

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# 1 Purpose

The purpose of this release note is to provide a brief description of the Dispense Record Structured Document Template specification for Electronic Transfer of Prescription Release 1 (ETP R1). This release note details the location of the specification, highlights resolved and outstanding issues, and also provides links to supporting documentation.

## 2 Introduction

The Dispense Record Structured Document Template for ETP R1 specifies the content of a prescribing exchange service dispense record (PES-DR) which is created by an electronic dispensing system (EDS) that is used by a community pharmacy. A PES-DR is created by an EDS as the record of dispensing of a prescribed item associated with an ePrescription. As such this specification complements the related ePrescription Structure Document Template specification (see section 4).

During dispensing, the dispenser supplies a therapeutic good, for the use by a subject of care (also known as the patient). As far as possible, the dispenser ensures that the subject of care receives the correct therapeutic good in the correct strength, dose form and quantity, and that the product is appropriately packed and labelled. Multiple dispenses (repeats) could take place against a single prescribed item and each is recorded as a PES-DR with reference to the source ePrescription.

The content of the specification will form the basis of mapping to the HL7 Clinical Document Architecture Version 2.0 (HL7 CDA V2) standard adopted in Australia for transmission of electronic clinical documents. These mappings will form the primary content of the Dispense Record CDA Implementation Guide for ETP R1. The specification can also, on a stakeholder needs basis, be used to map to other electronic message exchange formats such as HL7 v2.x.

The Dispense Record Structured Document Template for ETP R1 defines the value domains populated by code sets, specific concepts or reference sets built from SNOMED CT, SNOMED CT-AU or the Australian Medicines Terminology (AMT). For further information regarding terminology and the development of reference sets please visit <http://www.nehta.gov.au/connectingaustralia/clinical-terminologies> and direct your questions or feedback to [terminologies@nehta.gov.au](mailto:terminologies@nehta.gov.au).

NEHTA Clinical Information encourages feedback to this specification which can be submitted to [clinicalinfo@nehta.gov.au](mailto:clinicalinfo@nehta.gov.au).

## 3 Release Log

This specification is released with known issues that will be resolved in the next published version. Areas requiring resolution are recorded in the APPENDIX A: KNOWN ISSUES section of the specification.

## 4 Release Documentation

The following supporting documentation is available for download from the [NEHTA website](#).

[ETP Dispense Record - Structured Document Template - Release 1 v1.0 DRAFT](#)

[ETP ePrescription - Structured Document Template - Release 1 v1.0 DRAFT](#)

[ETP Business Process and Requirements Specification – Release 1 v1.0 DRAFT](#)

[ETP Technical Requirements Specification – Release 1 Release 1 v1.0 DRAFT](#)

[ETP Logical Information Model – Release 1 v1.0 DRAFT](#)

[ETP Technical Architecture – Release 1 v1.0 DRAFT](#)

[Data Specifications – Guide for Use v1.0](#)

[Document Header Data Specification v2.0](#)

[Participation Data Specification v2.0](#)

[Participation Specialisations Data Specification v1.0](#)

### 4.1 Dispense Record - Structured Document Template

The Dispense Record Structured Document for ETP R1 specifies the required data elements of a PES-DR, this specification includes their organisation (grouping), definition, datatype and constraints (including occurrence frequency, value domain and conditions of use).

#### File Details:

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