AUSTRALIAN MEDICINES TERMINOLOGY (AMT) DESCRIPTIONS for PHARMACEUTICAL BENEFITS SCHEME LISTINGS

The AMT is a standards-based national approach to the unique identification (code) and naming of medicines. It has been primarily developed to be used in clinical software applications and to facilitate interoperability between systems. For more information on the AMT and other Australian Digital Health Agency (the Agency) clinical terminology products and services, please visit the National Clinical Terminology Service website at https://www.healthterminologies.gov.au/

# SPONSOR TO COMPLETE

## STEP 1 - Sponsor to provide the NCTS team the requested information for description creation.

### Sponsor:

### Medicine (ingredient/s):

### Brand:

|  | Product Information(More than one formulation/strength/pack size can be entered for the same drug) |
| --- | --- |
| Trade name  | (The full exact name of the product as it appears on the label. Please replicate casing and punctuation exactly as it appears on the label) |
| Ingredients and strengths  | (The full detailed name of the ingredients and its associated strength in the product) |
| Dose form | (The form of the manufactured product) |
| PBS pack size  | (The total number of units contained within the manufacturer’s packaging for PBS listing) |
| Registered pack sizes  | (The total number of units contained within the manufacturer’s packaging for marketing in Australia) |
| Container type  | (The container immediately around the unit of use) |
| ARTG ID  | (Australian Register of Therapeutic Goods ID) |

## Sponsor contact details (for NCTS enquiries)

### Name of representative:

### Email:

### Phone:

### Due date for receipt of NCTS response (please allow at least 3 to 5 working days):

## STEP 2 - Sponsor to attach ARTG registration details, Product Label, Product Information and Consumer Medicines Information documents (where available) to ensure descriptions are aligned, accurate and unambiguous. These documents will not be distributed outside of the NCTS team.

### a) Please indicate below the type of information provided.

[x]  Product Label

[x]  Product Information (PI)

[x]  Consumer Medicines Information (CMI)

[x]  ARTG registration details

### b) Email to: help@digitalhealth.gov.au

# NCTS TEAM TO COMPLETE

## STEP 3 - NCTS team to complete this concept description table and send the completed form back to the sponsor (please copy and create more than one table for input if multiple concepts are involved).

|  |  |  |  |
| --- | --- | --- | --- |
| Concept SCT-ID\* | Concept Class | Reference Set Membership | Preferred Term Description |
|  | Substance (Basis of Strength Substance –BoSS)\*\* |  |  |
|  | Medicinal product (MP) | Medicinal Product (MP) |  |
|  | Clinical drug (CD) | Medicinal Product Unit of Use (MPUU) |  |
|  | Clinical drug package (CDP) | Medicinal Product Pack (MPP) |  |
|  | Product name (PN) | Trade Product (TP) |  |
|  | Branded clinical drug (BCD) | Trade Product Unit of Use (TPUU) |  |
|  | Branded clinical drug package (BCDP) | Trade Product Pack (TPP) |  |
|  | Containerized branded clinical drug package (CBCDP) | Containered Trade Product Pack (CTPP) |  |

Note:

\* Concept SCT-ID may not be provided in all circumstances

\*\* The BoSS is the name of the ingredient that the strength of the product is based on and has no associated SCT-ID.

**NCTS contact details for any sponsor enquiries**

**Service Desk ticket number:**

**Name of NCTS representative:**

**Email:** help@digitalhealth.gov.au

**Phone:** 1300 901 001

**Date of reply to sponsor:**

**Additional comments to sponsor (optional):**

# SPONSOR TO COMPLETE

## STEP 4 - Sponsor to provide completed form to PBAC Secretariat with submission.