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|  | **NHS Prescription Services**Bridge House152 Pilgrim StreetNewcastle upon TyneNE1 6SN0845 850 0001 Email:nhsbsa.prescriptionservices@nhsbsa.nhs.ukWebsite:[www.nhsbsa.nhs.uk](http://www.nhsbsa.nhs.uk)  24 Feb 2025 |
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VERSION 2.3.0

Dear dm+d User,

**Updates to concepts associated with VMP Varenicline 1mg tablets and Varenicline 500microgram tablets**

The dm+d extract files last week contained a duplicate 53 tablet VMPP (albeit made up of different components).

Authoring of dm+d has been updated in line with the table (see Invalidation table below) detailing concept descriptions and associated SNOMED codes.

In summary:

* The 53 tablet VMPP authored 10/02/2025 has been made invalid.
* The historical 53 tablet VMPP, which consisted of 3 components (11 x 0.5mg tablets, 14 x 1mg tablets and 28 x 1mg tablets), has been updated to 11 x 0.5mg tablets and 42 x 1mg tablets (newly authored) components.
* At the AMPP level, the AMPP authored 10/02/2025 has been made invalid.
* A new 42 tablet AMPP component has been authored linked to Accord UK Ltd
* For the discontinued AMP Champix 0.5mg/1mg 4 week treatment initiation pack (Pfizer Ltd) the AMPP level information has been updated with the 11 tablet component and linking made to the newly authored 42 tablet component.

The above changes allow varenicline initiation packs with the same components to share the same VMPP, regardless of packaging differences.

**Imdylltra powder for concentrate and solution for solution for infusion vials (all strengths)**

Due to product launch delay, the dm+d records for Imdylltra® (Tarlatamab, strengths: 1mg and 10mg) will temporarily be marked as not available until these products are marketed.

**Update on impact of reuse of identifiers**

**We are extending the deadline to seek further feedback from users**. We would appreciate feedback preferably no later than **Friday 7th March**, although suppliers are welcome to email us at dmdenquiries@nhsbsa.nhs.uk with any comments, concerns or queries beyond this date.

The recent work on adding ingredients to Virtual Therapeutic Moieties (VTMs) has been carried out. Where the VTM identifier is an International SNOMED identifier for a “contains only” Medicinal product concept then the assigned ingredient concept must also have an International SNOMED identifier.

This work has identified an issue.

* Currently the dm+d Editorial Policy states that SNOMED identifiers must not be re-used or assigned to another concept even where the concept identifier has been subsequently updated in dm+d.
* Appendix XXVI of the dm+d Editorial Policy details five ingredient concepts that have a SNOMED CT UK Extension identifier in dm+d but they have an International identifier in the SNOMED CT UK Drug Extension. These ingredient identifiers cannot be updated to International identifiers
* Because of the above points the VTM concepts that should have these ingredient concepts assigned to them do not currently have ingredients in dm+d.
* Appendix XXVI is in the process of being updated to include another ingredient concept where the international identifier cannot be allocated – Perflutren.
* The SNOMED CT UK Drug Extension handles the above through the use of both current and historical, relationship types for example ‘replaced by’, ‘same as’ etc. Also inactive and active concepts are identified in SNOMED but dm+d is unable to do this for ingredient concepts.
* Currently in dm+d if these ingredient concepts were required to be assigned to the VTM concepts it would only be possible to create the ingredient concept with a UK Extension identifier in dm+d. This UK Extension identifier would be inactive in the SNOMED CT UK Drug Extension with a SAME\_AS relationship to the International identifier concept
* This therefore introduces misalignment between the same VTM concepts in dm+d and the SNOMED CT UK Drug Extension.

Concern has been raised that as the terminologies evolve, more examples of this misalignment between them will come to light over time. Prior to proceeding with the assignation (or not) of ingredients to these VTMs in dm+d, the dm+d and NHSE Terminology teams sought feedback from system suppliers on the potential impact of reusing these particular codes (note – any reuse would always be noted in outgoing comms to alert system suppliers).

For the example below, suppliers were asked, “*what would be the impact upon your system and users* ***IF*** *the ingredient Sevelamer (or any of the other examples in the appendix) was assigned the international ID*”.

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| **Ingredient** | **dm+d concept ID** | **Status** |
| Sevelamer hydrochloride | 395871000 Sevelamer (substance) International IDFrom 08/2005 until 12/2010 | History – previous ID |
| 18308811000001104 - UK extension ID From 12/2010 to date | Current |
| Sevelamer | 18308711000001107 – UK extension IDFrom 12/2010 to date | Current |

In SNOMED UK Drug Extension:

395871000 Sevelamer (substance) – Active

18308811000001104  - Inactive

18308711000001107 - Inactive

The correct international identifier for the ingredient Sevelamer is 395871000 but this cannot be assigned as it was used between 2005 and 2010 and is part of the historical information for the ingredient Sevelamer hydrochloride.

**SNOMED International Proposal to Increase Description Length Limit**

We are providing an update to the [information previously communicated](https://nhsengland.kahootz.com/t_c_home/viewBlogArticle?articleID=1108441&nextURL=%2Ft_c_home%2FviewBlog%3Fblogid%3D50136) regarding the proposal from SNOMED International to increase the maximum length of Fully Specified Name and Synonym descriptions,.

SNOMED International have provided [further information](https://nhsengland.kahootz.com/t_c_home/viewBlogArticle?articleID=1154361&nextURL=%2Ft_c_home%2FviewBlog%3Fblogid%3D50136) about the [proposal to increase description length limit](https://nhsengland.kahootz.com/t_c_home/viewBlogArticle?articleID=1108441&nextURL=%2Ft_c_home%2FviewBlog%3Fblogid%3D50136) from 255 to 4096 characters.

NHS England are currently assessing whether this proposed change will impact Fully Specified Names (FSN) created in the SNOMED CT UK Drug Extension. NHS England do not expect that this proposed change will impact Preferred Terms (PT) in the SNOMED CT UK Drug Extension.

This proposal will impact UK users of the **Medicinal Product Hierarchy of the SNOMED International Edition** (both SNOMED CT International Edition and SNOMED CT UK Edition) but dm+d will NOT be affected and we have no plans to adopt these changes.

The planned implementation date for **SNOMED International** **changes is January 2026** and will come into effect in the **April 2026 UK release**.

Any concerns or queries should be directed to SNOMED International at release@snomed.org. Please cc information.standards@nhs.net to enable NHS England to have visibility of UK stakeholder feedback.

**Addition of 'Ingredients for VTMs' in dm+d (XML) release files and SNOMED CT UK Drug Extension**

The recent work to add the ingredients to Virtual Therapeutic Moieties (VTMs) that needed populating has been completed.

The [dm+d Editorial Policy](https://www.nhsbsa.nhs.uk/pharmacies-gp-practices-and-appliance-contractors/dictionary-medicines-and-devices-dmd) will be updated shortly to reflect the latest policy and will continue to be updated as-and-when any other exclusions are identified. We will provide confirmation when this has been completed.

**Please note that VTM Ingredients information will only be available via TRUD and the NHS England Terminology Server from 23 September 2024 and will not be seen in the dm+d browser until further notice is provided.**

Additional information about the change is available on the following [webpage](https://digital.nhs.uk/services/terminology-and-classifications/uk-medicines-terminology-futures/changes-to-digital-terminologies/addition-of-ingredients-for-vtms-in-dm-d-xml-release-files-and-snomed-ct-uk-drug-extension).

If you have any specific queries relating to the dm+d and SNOMED CT UK Drug Extension changes, please contact nhsdigital.ukmeds@nhs.net

**Monthly dm+d Supplier Workshop**

NHS England hosts a monthly workshop for IT system suppliers to discuss dm+d content and implementation queries.  It is an opportunity to speak with colleagues from NHS England and the NHSBSA who manage the terminology.

If you are a system supplier who is interested in joining the group, please contact medicinestandards@nhs.net for more information.

**DHSC require these concepts to be made available and visible (similar to licensed medicines) in prescribing and dispensing systems for the time being.**

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| **Specials/Imports added at request of DHSC to mitigate shortages in the supply chain** |
| **AMP Name** | **AMP SNOMED Code** | **Import/Special** | **dm+d extract date** |
| Desmopressin 150micrograms/dose nasal spray | 38955011000001100 | Imported  |  14/09/2020 |
| Desmopressin 150micrograms/dose nasal spray | 38996011000001101 | Special Order |  21/09/2020 |
| Capsaicin 0.025% cream | 41539411000001105 | Imported |  27/03/2023 |
| Capsaicin 0.075% cream | 41540711000001105 | Imported |  27/03/2023 |
| Griseofulvin 125mg tablets | 42441511000001104 | Imported (United States) |  27/11/2023 |
| Griseofulvin 500mg tablets | 42441811000001101 | Imported (United States) |  27/11/2023 |
| Hydrogen peroxide 3% solution  | 42510811000001109 | Imported |  18/12/2023 |
| Cerium nitrate hexahydrate 22mg/g / Sulfadiazine silver 10mg/g cream | 42991011000001101 | Imported |  01/07/2024 |
| Sulfadiazine silver 1% cream  | 42990711000001107 | Imported | 01/07/2024 |
| Creon 10000 capsules | 43034611000001101 | Imported | 15/07/2024 |
| Creon 25000 capsules | 43034811000001102 | Imported | 15/07/2024 |
| Pancreaze Delayed-Release capsules | 43035111000001108 | Imported | 15/07/2024 |
| Ipratropium bromide 250micrograms/1ml nebuliser liquid unit dose vials | 43093511000001105 | Imported | 22/07/2024 |
| Ipratropium bromide 500micrograms/2ml nebuliser liquid unit dose vials | 43094411000001109 | Imported | 22/07/2024 |
| Methylphenidate 18mg modified-release tablets | 43210911000001100 | Imported  | 02/09/2024 |
| Methylphenidate 27mg modified-release tablets | 43211511000001100 | Imported | 02/09/2024 |
| Methylphenidate 36mg modified-release tablets | 43211911000001107 | Imported | 02/09/2024 |
| Methylphenidate 54mg modified-release tablets | 43212411000001109 | Imported | 02/09/2024 |
| Pangrol 10,000 capsules | 43788211000001100 | Imported | 23/09/2024 |
| Pangrol 25,000 capsules | 43788511000001102 | Imported | 23/09/2024 |
| Imiquimod 5% cream 250mg sachets | 41526711000001106 | Imported | 02/12/2024 |
| Quetiapine 50mg modified-release tablets | 44616911000001103 | Imported | 17/02/2025 |
| Quetiapine 150mg modified-release tablets | 44617411000001108 | Imported | 17/02/2025 |
| Quetiapine 200mg modified-release tablets | 44617711000001102 | Imported | 17/02/2025 |
| Quetiapine 300mg modified-release tablets | 44618011000001103 | Imported | 17/02/2025 |
| Quetiapine 400mg modified-release tablets | 44618311000001100 | Imported | 17/02/2025 |
| Fluorouracil 5% cream | 44618611000001105 | Imported | 17/02/2025 |
| Imiquimod 3.75% cream 250mg sachets | 44618911000001104 | Imported | 17/02/2025 |
| Sodium fusidate 250mg tablets | 44619211000001103 | Imported | 17/02/2025 |
| Fusidic acid 250mg/5ml oral suspension | 44619511000001100 | Special Order | 17/02/2025 |

**Invalidation****s**

The following concept has been invalidated (see above).

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| **Concept to be made Invalid** | **Replacement concepts** |
| **VMP**Varenicline 1mg tablets and Varenicline 500microgram tablets (Not to be made invalid) | **VMP**N/A |
| **VMP SNOMED ID**10984311000001109  | **VMP SNOMED ID**N/A |
| **VMPP**53 tablets | **VMPP**N/A |
| **VMPP SNOMED ID**44562711000001106 | **VMPP SNOMED ID**N/A |

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| **Concept to be made Invalid** | **Replacement concepts** |
| **VMP**Varenicline 1mg tablets and Varenicline 500microgram tablets(Not to be made invalid) | **VMP**N/A |
| **VMP SNOMED ID**10984311000001109  | **VMP SNOMED ID**N/A |
| **VMPP**53 tablets (Not to be made invalid) | **VMPP**N/A |
| **VMPP SNOMED ID**44562711000001106  | **VMPP SNOMED ID**N/A |
| **AMP**Varenicline 1mg tablets and Varenicline 500microgram tablets (Accord-UK Ltd)(Not to be made invalid) | **AMP**N/A |
| **AMP SNOMED ID**44551411000001101 | **AMP SNOMED ID**N/A |
| **AMPP**53 tablets | **AMPP**53 tablets |
| **AMPP SNOMED ID**44562811000001103 | **AMPP SNOMED ID**44742211000001108 |

**Advance Notice of Invalidations**

The following concept will be invalidated mid-March as it was added erroneously.

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| **Concept to be made Invalid** | **Replacement concepts** |
| **VMP**Amikacin 500mg/2ml solution for injection vials (not to be made invalid) | **VMP**Amikacin 500mg/2ml solution for injection ampoules |
| **VMP SNOMED ID**35899911000001109 | **VMP SNOMED ID**44422511000001103 |
| **VMPP**5 vials (not to be made invalid) | **VMPP**5 ampoule |
| **VMPP SNOMED ID**4500011000001105  | **VMPP SNOMED ID**44374911000001108  |
| **AMP**Amikacin 500mg/2ml solution for injection vials (Kent Pharma (UK) Ltd) | **AMP**Amikacin 500mg/2ml solution for injection ampoules (Kent Pharma (UK) Ltd) |
| **AMP SNOMED ID** 42788511000001107  | **AMP SNOMED ID**44375011000001108 |
| **AMPP**5 vials | **AMPP**5 ampoule |
| **AMPP SNOMED ID**42788611000001106 | **AMPP SNOMED ID**44375111000001109  |

dm+d Authoring Team