# **Data Changes for the CCDD DECEMBER Release Candidate 2021**

## **Based on the QA Release Generation** 20211201

|  |  |
| --- | --- |
| Data entered by | J. Hutsul |
| Data checked by | A. Raghuveer |

## **TM File**

### **Deprecation**

The following TM concept(s) needs to be added back into the TM File with a status of **deprecated**. [Reason for deprecation].

|  |  |  |
| --- | --- | --- |
| **tm\_code** | **tm \_formal\_name** | **tm\_status** |
|  | [NONE] | Deprec |

### **Code permanence – name changes**

The following TM concept(s) need to keep its/their existing tm\_code even though their formal name has changed. [Reason for name change].

|  |  |  |
| --- | --- | --- |
| **tm\_code (to keep)** | **tm \_formal\_name** | **tm\_code in QA Release was** |
|  | [NONE] |  |

### **Concept permanence - returns**

The following TM concept(s) is/are “missing” from the QA Release, but should come back ‘automatically’ in the Release Candidate with a status of “inactive”, when its/their supporting NTPs and MPs will be on the “Whitelist”, as their DINs have been Cancelled Post Market or are Dormant.

|  |  |
| --- | --- |
| **tm\_code** | **tm \_formal\_name** |
| 8001835 | levobunolol |

## **NTP File**

### **Deprecation**

The following NTP concept(s) need to be added back into the NTP File with status of **deprecated**.

|  |  |  |  |
| --- | --- | --- | --- |
| **ntp\_code** | **ntp \_formal\_name** | **ntp\_fr\_description** | **ntp\_status** |
|  | [NONE] |  |  |

### **Code permanence – name changes**

The following NTP concepts need to keep their existing ntp\_code even though their formal name has changed.

|  |  |  |
| --- | --- | --- |
| **ntp\_code (to keep)** | **ntp \_formal\_name** | **ntp\_code in QA Release was** |
| 9014447 | buprenorphine (buprenorphine hydrochloride) 12 mg and naloxone (naloxone hydrochloride) 3 mg oromucosal film | 0ac5384c89524f7642d78537e24ea00b |
| 9014448 | buprenorphine (buprenorphine hydrochloride) 2 mg and naloxone (naloxone hydrochloride) 0.5 mg oromucosal film | 369807a16f501b9798077d37b9b7c905 |
| 9014541 | buprenorphine (buprenorphine hydrochloride) 4 mg and naloxone (naloxone hydrochloride) 1 mg oromucosal film | f5128bcac35e90e593399c081af55197 |
| 9014449 | buprenorphine (buprenorphine hydrochloride) 8 mg and naloxone (naloxone hydrochloride) 2 mg oromucosal film | ab91cda0b7fbd9bcbfe03a8431ac2733 |

### **Concept permanence – returns check**

The following NTP concepts are “missing” from the QA Release, but should come back ‘automatically’ in the Release Candidate, when their supporting MPs will be on the “Whitelist”, as their DINs have been Cancelled Post Market or are Dormant.

|  |  |
| --- | --- |
| **ntp\_code** | **ntp \_formal\_name** |
| 9001940 | dipyridamole 25 mg oral tablet |
| 9001941 | dipyridamole 50 mg oral tablet |
| 9002836 | levobunolol hydrochloride 0.5 % ophthalmic drops |
| 9001677 | methylprednisolone (methylprednisolone sodium succinate) 1 g per vial powder for solution for injection |
| 9003470 | methylprednisolone (methylprednisolone sodium succinate) 40 mg per vial powder for solution for injection |
| 9001664 | methylprednisolone (methylprednisolone sodium succinate) 500 mg per vial powder for solution for injection |

## **MP File**

### **Deprecation (very rare)**

No new deprecated concepts.

### **Deprecation – manual return**

The following MP concepts need to be added back into the MP File with status of **deprecated**. For all DINs, they have had additional UoP added. All products now have new MP.

|  |  |  |  |
| --- | --- | --- | --- |
| **mp\_code** | **mp \_formal\_name** | **mp\_fr\_description** | **mp\_status** |
| 00587737 | HUMULIN N (insulin isophane human 100 unit per mL suspension for injection 10 mL vial) ELI LILLY CANADA INC | HUMULIN N (insuline isophane humaine 100 unité par mL suspension injectable 10 mL fiole) ELI LILLY CANADA INC | Deprec |
| 02182971 | METHOTREXATE INJECTION USP (methotrexate (methotrexate sodium) 1000 mg per 40 mL solution for injection vial) PFIZER CANADA ULC | METHOTREXATE INJECTION USP (méthotrexate (méthotrexate sodique) 1000 mg par 40 mL solution injectable fiole) PFIZER CANADA ULC | Deprec |
| 02182777 | METHOTREXATE INJECTION USP (methotrexate (methotrexate sodium) 500 mg per 20 mL solution for injection vial) PFIZER CANADA ULC | METHOTREXATE INJECTION USP (méthotrexate (méthotrexate sodique) 500 mg par 20 mL solution injectable fiole) PFIZER CANADA ULC | Deprec |

The following MP concepts are generating as active but need to be set to **deprecated**. For both products, they have been assigned a new DIN (02474891, 02474735). This will have to be done manually each month.

|  |  |  |
| --- | --- | --- |
| **mp\_code** | **mp \_formal\_name** | **mp\_status** |
| 77700322 | METHOTREXATE INJECTION BP (methotrexate (methotrexate sodium) 1000 mg per 40 mL solution for injection vial) ACCORD HEALTHCARE INC | Deprec |
| 77700323 | METHOTREXATE INJECTION BP (methotrexate (methotrexate sodium) 500 mg per 20 mL solution for injection vial) ACCORD HEALTHCARE INC | Deprec |
| 77700627 | DIMENHYDRINATE INJECTION USP (dimenhydrinate 250 mg per 5 mL solution for injection vial) TELIP, LLC, A SUBSIDIARY OF TELIGENT, INC. | Deprec |

### **DPD descriptors**

The following concept(s) needs to have the DPD descriptor added to the brand name to allow differentiation between duplicate mp\_formal names.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| drug\_  code | mp\_code | mp\_formal\_name | DPD descriptor | DPD descriptor - FR |
|  | [NONE] |  |  |  |

### **Concept permanence – manual return**

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| drug\_code | mp\_code | mp\_formal\_name | mp\_fr\_description | CCDD status |
|  | [NONE] |  |  |  |

### **Concept permanence – returns check**

The following MPs need to be added to the Whitelist to make sure that they (and their associated NTPs/TMs) continue to be present in the release:

|  |  |  |  |
| --- | --- | --- | --- |
| **drug\_code** | **mp\_code** | **mp\_formal\_name** | **DPD status** |
| 90350 | 2418959 | ALLERTIN (loratadine 10 mg oral tablet) APOTEX INC | Dormant 2021.10.14 |
| 11034 | 895644 | APO-DIPYRIDAMOLE (dipyridamole 25 mg oral tablet) APOTEX INC | Dormant 2021.09.25 |
| 11035 | 895652 | APO-DIPYRIDAMOLE (dipyridamole 50 mg oral tablet) APOTEX INC | Dormant 2021.09.25 |
| 78557 | 2300257 | BALMINIL DECONGEST (xylometazoline hydrochloride 0.1 % nasal spray) TEVA CANADA LIMITED | cancelled post-market Expiry 2021.11.30 |
| 6230 | 637661 | BETAGAN 0.5% OPHTHALMIC SOL (levobunolol hydrochloride 0.5 % ophthalmic drops) ALLERGAN INC | cancelled post-market Expiry 2021.11.30 |
| 81075 | 2325357 | BUPROPION SR (bupropion hydrochloride 150 mg prolonged-release oral tablet) PRO DOC LIMITEE | cancelled post-market Expiry 2021.11.30 |
| 83716 | 2351927 | COLD AND ALLERGY DECONGESTANT NASAL SPRAY (xylometazoline hydrochloride 0.1 % nasal spray) TEVA CANADA LIMITED | cancelled post-market Expiry 2021.11.30 |
| 69738 | 2245638 | CORICIDIN COLD & FLU EXTRA STRENGTH (acetaminophen 500 mg and chlorpheniramine maleate 2 mg and dextromethorphan hydrobromide 15 mg oral tablet) BAYER INC | cancelled post-market Expiry 2021.11.30 |
| 41933 | 1919245 | DR. SCHOLL'S ATHLETE'S FOOT SPRAY POWDER (tolnaftate 1 % cutaneous spray powder) SCHOLL'S WELLNESS COMPANY | Dormant 2021.10.29 |
| 97070 | 2480166 | FUNGIFOAM ANTIFUNGAL (tolnaftate 1 % cutaneous cream) THE TETRA CORPORATION | cancelled post-market Expiry 2020.12.31 |
| 18442 | 2158442 | HYGENIPAK GREEN ANTI-MICROBIAL SKIN CLEANSER (chloroxylenol 0.15 % and triclosan 0.15 % cutaneous gel) DEB CANADA | cancelled post-market Expiry 2021.11.30 |
| 88567 | 2400871 | KETOTIFEN OPHTHALMIC SOLUTION (ketotifen (ketotifen fumarate) 0.25 mg per mL ophthalmic drops) STERIMAX INC | Dormant 2021.08.19 |
| 94136 | 2454653 | LEVETIRACETAM (levetiracetam 250 mg oral tablet) PHARMASCIENCE INC | Dormant 2021.03.03 |
| 94137 | 2454661 | LEVETIRACETAM (levetiracetam 500 mg oral tablet) PHARMASCIENCE INC | Dormant 2020.09.19 |
| 94138 | 2454688 | LEVETIRACETAM (levetiracetam 750 mg oral tablet) PHARMASCIENCE INC | Dormant 2021.03.04 |
| 88350 | 2398559 | LIDOCAINE HYDROCHLORIDE SOLUTION (lidocaine hydrochloride 1 % cutaneous spray) PRIME ENTERPRISES INC | cancelled post-market Expiry 2021.11.30 |
| 87682 | 2391473 | MAR-GABAPENTIN (gabapentin 100 mg oral capsule) MARCAN PHARMACEUTICALS INC | Dormant 2020.12.18 |
| 87683 | 2391481 | MAR-GABAPENTIN (gabapentin 300 mg oral capsule) MARCAN PHARMACEUTICALS INC | Dormant 2021.04.05 |
| 87684 | 2391503 | MAR-GABAPENTIN (gabapentin 400 mg oral capsule) MARCAN PHARMACEUTICALS INC | Dormant 2021.02.05 |
| 64837 | 2241229 | METHYLPREDNISOLONE SODIUM SUCCINATE FOR INJECTION (methylprednisolone (methylprednisolone sodium succinate) 1 g per vial powder for solution for injection) TEVA CANADA LIMITED | Dormant 2021.11.17 |
| 51346 | 2231893 | METHYLPREDNISOLONE SODIUM SUCCINATE FOR INJECTION (methylprednisolone (methylprednisolone sodium succinate) 40 mg per vial powder for solution for injection) TEVA CANADA LIMITED | Dormant 2021.11.17 |
| 51348 | 2231895 | METHYLPREDNISOLONE SODIUM SUCCINATE FOR INJECTION (methylprednisolone (methylprednisolone sodium succinate) 500 mg per vial powder for solution for injection) TEVA CANADA LIMITED | Dormant 2021.11.17 |
| 50886 | 2231543 | PMS-CARBAMAZEPINE-CR (carbamazepine 200 mg prolonged-release oral tablet) PHARMASCIENCE INC | Dormant 2021.03.05 |
| 50887 | 2231544 | PMS-CARBAMAZEPINE-CR (carbamazepine 400 mg prolonged-release oral tablet) PHARMASCIENCE INC | Dormant 2021.03.19 |
| 79685 | 2311267 | PRO-DEXAMETHASONE - 4 (dexamethasone 4 mg oral tablet) PRO DOC LIMITEE | cancelled post-market Expiry 2021.11.30 |
| 11300 | 1902660 | RETROVIR (AZT) (zidovudine 100 mg oral capsule) VIIV HEALTHCARE ULC | cancelled post-market Expiry 2021.11.30 |
| 66848 | 2243007 | SEEQUIN 4 IDS (hydroquinone 4 % cutaneous gel) VIVIER CANADA INCORPORATED | cancelled post-market Expiry 2021.12.01 |
| 98326 | 2493284 | TEA TREE OIL 2IN1 (pyrithione zinc 1 % shampoo) PROCTER & GAMBLE INC | cancelled post-market Expiry 2021.11.20 |
| 89497 | 2410338 | TETRABENAZINE TABLETS (tetrabenazine 25 mg oral tablet) STERIMAX INC | Dormant 2021.10.21 |
| 91438 | 2429969 | TRAMADOL / ACETAMINOPHEN (acetaminophen 325 mg and tramadol hydrochloride 37.5 mg oral tablet) SIVEM PHARMACEUTICALS ULC | cancelled post-market Expiry 2021.11.09 |
| 6462 | 632732 | VOLTAREN (diclofenac sodium 100 mg suppository) NOVARTIS PHARMACEUTICALS CANADA INC | cancelled post-market Expiry 2017.11.30 |
| 4043 | 514012 | VOLTAREN (diclofenac sodium 50 mg gastro-resistant tablet) NOVARTIS PHARMACEUTICALS CANADA INC | cancelled post-market Expiry 2020.08.31 |
| 9943 | 782459 | VOLTAREN SR (diclofenac sodium 75 mg prolonged-release oral tablet) NOVARTIS PHARMACEUTICALS CANADA INC | cancelled post-market Expiry 2020.05.31 |
| 100004 | 2510847 | VAXZEVRIA (ChAdOx1-S (recombinant) 5e10 vp per 0.5 mL solution for injection 5 mL vial) ASTRAZENECA CANADA INC | Approved through regular submission process |

## **Blacklist (on GitHub)**

The following changes have been made to the Blacklist for the August 2021 Release Candidate. The list as it is on GitHub should be used.

Changes that have been made to the Blacklist are as follows:  
ACTION is either:

* STAY on Blacklist
* ADD to Blacklist
* REMOVE from Blacklist

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **ntp\_code** | **ntp\_formal\_name** | **drug\_code** | **mp\_code** | **mp\_formal\_name** | **ACTION** |
|  | lidocaine 0.8 % cutaneous gel | 98886 | 02499118 | AFTER SUN GEL (lidocaine 0.8 % cutaneous gel) FRUIT OF THE EARTH, INC | STAY |
| 9014551 | methotrexate (methotrexate disodium) 2.5 mg oral tablet | 99838 | 02509067 | ACH-METHOTREXATE (methotrexate (methotrexate disodium) 2.5 mg oral tablet) ACCORD HEALTHCARE INC | REMOVE |