# Data Changes for the CCDD APR Release Candidate 2020

## Based on the QA Release 20200401

|  |  |
| --- | --- |
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## TM File

### Deprecation

The following 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 concept(s) need to keep 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

In addition, the following concept is “missing” from the QA Release, but should come back ‘automatically’ in the Release Candidate with a status of “inactive”, when its 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** |
| 8001509 | polidocanol |

## NTP File

### Deprecation

The following concept(s) need to be added back into the NTP File with status of **deprecated**. The ntp\_formal\_name is longer being used since the dosage form of the only product associated with it changed in DPD from powder for solution to granules for suspension.

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

### Code permanence – name changes

The following 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** |
|  | [NONE] |  |

### Concept permanence – returns check

The following four 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** |
| 9013132 | follitropin alfa 450 unit per vial powder for solution for injection with diluent solution syringe |
| 9009882 | hydrocodone bitartrate 5 mg oral tablet |
| 9000278 | metoprolol tartrate 5 mg per 5 mL solution for injection ampoule |
| 9000674 | metronidazole 0.75 % cutaneous cream |
| 9013468 | ocriplasmin 1.25 mg per mL solution for injection 0.3 mL vial |
| 9003831 | petrolatum 49.5 % and zinc oxide 40 % cutaneous ointment |
| 9013835 | polidocanol 1.3 mg per mL injectable foam 45 mL canister |
| 9013247 | white petrolatum 99.9 % cutaneous ointment |

## MP File

### Deprecation (very rare)

**No new deprecated concepts**

### Deprecation – manual return

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

|  |  |  |  |
| --- | --- | --- | --- |
| **mp\_code** | **mp \_formal\_name** | **mp\_fr\_description** | **mp\_status** |
| 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 concepts are generating as active but need to be set to **deprecated**. For both products, they have been assigned a new DIN (02474891, 02474735). I believe 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 |

### DPD descriptors

The following concepts 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

One concept needs to be added back into the generation with its status as **active**. For this product, in the DPD it has gone back to Approved while the company “reactivates” its DIN.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| drug\_code | mp\_code | mp\_formal\_name | mp\_fr\_description | CCDD status |
| 15207 | 01916947 | BACTROBAN (mupirocin 2 % cutaneous ointment) GLAXOSMITHKLINE CONSUMER HEALTHCARE INC. | BACTROBAN (mupirocine 2 % pommade cutanée) GLAXOSMITHKLINE CONSUMER HEALTHCARE INC. | Active |

### 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 |
| 95194 | 02463849 | ACCEL-CANDESARTAN/HCTZ (candesartan cilexetil 32 mg and hydrochlorothiazide 12.5 mg oral tablet) ACCEL PHARMA INC | Cancelled post-market 2020-03-05 |
| 95195 | 02463857 | ACCEL-CANDESARTAN/HCTZ (candesartan cilexetil 32 mg and hydrochlorothiazide 25 mg oral tablet) ACCEL PHARMA INC | Cancelled post-market 2020-03-05 |
| 77281 | 02287404 | ACT SERTRALINE (sertraline (sertraline hydrochloride) 50 mg oral capsule) ACTAVIS PHARMA COMPANY | Cancelled post-market Expiry 2020-03-31 |
| 72398 | 02248076 | DIAPER RASH OINTMENT/ONGUENT POUR LE TRAITEMENT DE L'ÉRYTHÈME FESSIER (petrolatum 49.5 % and zinc oxide 40 % cutaneous ointment) LES PRODUITS DE SOINS POUR LA PEAU AU LAIT DE CHEVRE CANUS INC/CANUS GOAT'S MILK | Cancelled post-market Expiry 2020-03-24 |
| 88074 | 02395665 | FENTANYL PATCH (fentanyl 25 mcg per hour transdermal patch) PRO DOC LIMITEE | Cancelled post-market Expiry 2020-03-31 |
| 88075 | 02395673 | FENTANYL PATCH (fentanyl 50 mcg per hour transdermal patch) PRO DOC LIMITEE | Cancelled post-market Expiry 2020-03-31 |
| 88076 | 02395681 | FENTANYL PATCH (fentanyl 75 mcg per hour transdermal patch) PRO DOC LIMITEE | Cancelled post-market Expiry 2020-03-31 |
| 75998 | 02274531 | GD-AZITHROMYCIN (azithromycin 250 mg oral tablet) GENMED A DIVISION OF PFIZER CANADA ULC | Cancelled post-market Expiry 2020-03-31 |
| 72488 | 02248156 | GONAL-F (follitropin alfa 450 unit per vial powder for solution for injection with diluent solution syringe) EMD SERONO A DIVISION OF EMD INC CANADA | Cancelled post-market Expiry 2020-02-29 |
| 13314 | 01916599 | HYCODAN TABLETS (hydrocodone bitartrate 5 mg oral tablet) BRISTOL-MYERS SQUIBB CANADA | Cancelled post-market Expiry 2020-03-31 |
| 93860 | 02452154 | JETREA (ocriplasmin 1.25 mg per mL solution for injection 0.3 mL vial) THROMBOGENICS NV | Cancelled post-market 2020-03-20 |
| 7357 | 00590819 | LOPRESOR INJ 1.0MG/ML (metoprolol tartrate 5 mg per 5 mL solution for injection ampoule) NOVARTIS PHARMACEUTICALS CANADA INC | Cancelled post-market Expiry 2020-03-05 |
| 60162 | 02236872 | METHOXACET-C 1/8 (acetaminophen 325 mg and codeine phosphate 8 mg and methocarbamol 400 mg oral tablet) TEVA CANADA LIMITED | Cancelled post-market Expiry 2020-03-31 |
| 92647 | 02442116 | METOPROLOL-L (metoprolol tartrate 25 mg oral tablet) SIVEM PHARMACEUTICALS ULC | Cancelled post-market 2019-07-17 |
| 43445 | 02226839 | METROCREAM - CRM 0.75% (metronidazole 0.75 % cutaneous cream) GALDERMA CANADA INC | Cancelled post-market Expiry 2020-03-31 |
| 87825 | 02392992 | MINT-IRBESARTAN/HCTZ (hydrochlorothiazide 12.5 mg and irbesartan 150 mg oral tablet) MINT PHARMACEUTICALS INC | Dormant 2020-02-26 |
| 65251 | 02241594 | MIRAPEX (pramipexole dihydrochloride monohydrate 0.5 mg oral tablet) BOEHRINGER INGELHEIM (CANADA) LTD LTEE | Cancelled post-market Expiry 2020-03-31 |
| 60464 | 02237146 | MIRAPEX (pramipexole dihydrochloride monohydrate 1 mg oral tablet) BOEHRINGER INGELHEIM (CANADA) LTD LTEE | Cancelled post-market Expiry 2020-03-03 |
| 61247 | 02237886 | MYLAN-ACEBUTOLOL (TYPE S) (acebutolol (acebutolol hydrochloride) 200 mg oral tablet) MYLAN PHARMACEUTICALS ULC | Cancelled post-market Expiry 2020-03-31 |
| 64175 | 02240604 | MYLAN-AMIODARONE (amiodarone hydrochloride 200 mg oral tablet) MYLAN PHARMACEUTICALS ULC | Cancelled post-market Expiry 2020-03-31 |
| 79592 | 02310279 | MYLAN-VENLAFAXINE XR (venlafaxine (venlafaxine hydrochloride) 37.5 mg prolonged-release oral capsule) MYLAN PHARMACEUTICALS ULC | Cancelled post-market Expiry 2020-03-30 |
| 43291 | 02220636 | OXYBUTYNINE-5 - TAB 5MG (oxybutynin chloride 5 mg oral tablet) PRO DOC LIMITEE | Cancelled post-market Expiry 2020-03-31 |
| 9900 | 00828424 | PINDOLOL-10 TAB 10MG (pindolol 10 mg oral tablet) PRO DOC LIMITEE | Cancelled post-market 2020-03-09 |
| 9899 | 00828416 | PINDOLOL-5 TAB 5MG (pindolol 5 mg oral tablet) PRO DOC LIMITEE | Cancelled post-market 2020-03-09 |
| 93403 | 02448335 | SOLIFENACIN SUCCINATE TABLETS (solifenacin succinate 5 mg oral tablet) MDA INC. | Cancelled post-market 2020-03-06 |
| 92922 | 02444267 | VARITHENA (polidocanol 1.3 mg per mL injectable foam 45 mL canister) PROVENSIS LTD | Dormant 2020-02-28 |
| 80621 | 02320800 | WHITE PETROLEUM JELLY BABY FRESH SCENT (white petrolatum 99.9 % cutaneous ointment) BELVEDERE INTERNATIONAL INC. | Cancelled post-market Expiry 2020-03-31 |
| 80620 | 02320797 | WHITE PETROLEUM JELLY ULTRA MOISTURIZER (white petrolatum 100 % cutaneous ointment) BELVEDERE INTERNATIONAL INC. | Cancelled post-market Expiry 2020-03-31 |

## Blacklist (on GitHub)

Updates to the Blacklist to be applied for the DEC Release Candidate have been made. 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** |
| ecf2350d81789a8b037ee83efe8c33f3 | epinephrine 0.15 mg per 0.15 mL solution for injection | 86792 | 02382059 | ALLERJECT (epinephrine 0.15 mg per 0.15 mL solution for injection) KALEO INC | ADD |
| 592b2cb765438a31fa9163fb197b3319 | epinephrine 0.3 mg per 0.3 mL solution for injection | 86793 | 02382067 | ALLERJECT (epinephrine 0.3 mg per 0.3 mL solution for injection) KALEO INC | ADD |
| d20439bd8a022f1aa281716df5861c5a | sapropterin dihydrochloride 100 mg per sachet oral powder | 97268 | 2482207 | KUVAN (sapropterin dihydrochloride 100 mg per sachet oral powder) BIOMARIN INTERNATIONAL LIMITED | STAY |
| 48b077a2a72e4cc6a70cef04862e7c49 | sapropterin dihydrochloride 500 mg per sachet oral powder | 97269 | 2482215 | KUVAN (sapropterin dihydrochloride 500 mg per sachet oral powder) BIOMARIN INTERNATIONAL LIMITED | STAY |