# **Data Changes for the CCDD MARCH Release Candidate 2022**

## **Based on the QA Release Generation** 20220301

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

## **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] |  |

### **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** |
| 8002089 | aluminum chlorohydrate and methylprednisolone acetate and neomycin and sulfur |
| 8001751 | dimethicone and white petrolatum |

## **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** |
|  | [NONE] |  |

### **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** |
| 9001610 | aluminum chlorohydrate 100 mg per mL and methylprednisolone acetate 2.5 mg per mL and neomycin sulfate 2.5 mg per mL and sulfur 50 mg per mL lotion |
| 9005847 | dimethicone 10 % and white petrolatum 68.4 % cutaneous ointment |
| 9000230 | morphine sulfate 10 mg suppository |
| 9005846 | ponatinib (ponatinib hydrochloride) 45 mg oral tablet |

## **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 |
| 02258595 | HUMIRA (adalimumab 40 mg per 0.8 mL solution for injection vial) ABBVIE CORPORATION | HUMIRA (adalimumab 40 mg par 0,8 mL solution injectable fiole) ABBVIE CORPORATION | 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** |
| 9168 | 755885 | APO-PINDOL TAB 10MG (pindolol 10 mg oral tablet) APOTEX INC | Dormant 2022.01.20 |
| 64173 | 2240602 | BUSULFEX (busulfan 60 mg per 10 mL solution for injection vial) OTSUKA PHARMACEUTICAL CO LTD | cancelled post-market Expiry 2022.01.31 |
| 92798 | 2443333 | CAVILON MOISTURE BARRIER OINTMENT (dimethicone 10 % and white petrolatum 68.4 % cutaneous ointment) 3M CANADA COMPANY | Cancelled post-market 2022-02-28 |
| 70147 | 2246021 | COLD & ALLERGY RELIEF (xylometazoline hydrochloride 0.1 % nasal spray) PHARMASCIENCE INC | cancelled post-market Expiry 2022.02.28 |
| 83444 | 2349124 | EFFIENT (prasugrel (prasugrel hydrochloride) 10 mg oral tablet) ELI LILLY CANADA INC | cancelled post-market Expiry 2022.02.28 |
| 92169 | 2437341 | ICLUSIG (ponatinib (ponatinib hydrochloride) 45 mg oral tablet) ARIAD PHARMACEUTICALS INC | Dormant 2022.01.20 |
| 70148 | 2246022 | LONG LASTING NASAL MIST (oxymetazoline hydrochloride 0.05 % nasal spray) PHARMASCIENCE INC | cancelled post-market Expiry 2022.02.28 |
| 97248 | 2481979 | METHADONE HYDROCHLORIDE ORAL CONCENTRATE USP (methadone hydrochloride 10 mg per mL oral solution) STERINOVA INC. | cancelled post-market Expiry 2021.10.31 |
| 1394 | 195057 | NEO-MEDROL ACNE LOTION (aluminum chlorohydrate 100 mg per mL and methylprednisolone acetate 2.5 mg per mL and neomycin sulfate 2.5 mg per mL and sulfur 50 mg per mL lotion) PFIZER CANADA ULC | cancelled post-market Expiry 2022.02.28 |
| 78305 | 2297701 | NIGHTTIME SLEEP AID (diphenhydramine hydrochloride 25 mg oral capsule) APOTEX INC | Dormant 2022.01.23 |
| 50848 | 2231508 | PMS-DICLOFENAC (diclofenac sodium 100 mg suppository) PHARMASCIENCE INC | cancelled post-market Expiry 2022.01.31 |
| 50847 | 2231506 | PMS-DICLOFENAC (diclofenac sodium 50 mg suppository) PHARMASCIENCE INC | cancelled post-market Expiry 2022.02.28 |
| 50845 | 2231505 | PMS-DICLOFENAC-SR (diclofenac sodium 100 mg prolonged-release oral tablet) PHARMASCIENCE INC | cancelled post-market Expiry 2018.09.30 |
| 50844 | 2231504 | PMS-DICLOFENAC-SR (diclofenac sodium 75 mg prolonged-release oral tablet) PHARMASCIENCE INC | cancelled post-market Expiry 2022.01.31 |
| 6132 | 632201 | STATEX-SUPPOSITORIES 10MG (morphine sulfate 10 mg suppository) PALADIN LABS INC. | cancelled post-market Expiry 2022.02.28 |

## **Blacklist (on GitHub)**

The following changes have been made to the Blacklist for the March 2022 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 |