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

## **Based on the QA Release Generation** 20221003

|  |  |
| --- | --- |
| Data entered by | Jo-Anne Hutsul |
| Data checked by |  |

## **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** |
| 8001661 | hydroquinone |

## **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] |  |  |

### **Status update – Inactive**

The following NTP concept(s) are generating as Active but need to be set to **inactive** to align with their respective MPs.

|  |  |  |  |
| --- | --- | --- | --- |
| **ntp\_code** | **ntp \_formal\_name** | **ntp\_fr\_description** | **ntp\_status** |
| 9000881 | diazepam 2.5 mg per 0.5 mL rectal gel tube | diazépam 2,5 mg par 0,5 mL gel rectal tube | Inactive |
| 9000883 | diazepam 7.5 mg per 1.5 mL rectal gel tube | diazépam 7,5 mg par 1,5 mL gel rectal tube | Inactive |
| 9000880 | diazepam 20 mg per 4 mL rectal gel tube | diazépam 20 mg par 4 mL gel rectal tube | Inactive |

### **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** |
| 9012792 | diphenhydramine hydrochloride 25 mg and ibuprofen (ibuprofen, ibuprofen potassium) 200 mg oral capsule |
| 9013552 | exenatide 2 mg powder for prolonged-release suspension for injection with diluent solution per pen |
| 9013170 | interferon beta-1b 0.3 mg per vial powder for solution for injection with diluent solution syringe |

## **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 |

### **Status update – deprecation**

The following MP concepts are generating as active but need to be set to **deprecated**. Some of these products have been assigned a new DIN, others have been discontinued. 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 |
| 77700070 | DIASTAT (diazepam 15 mg per 3 mL rectal gel tube) BAUSCH HEALTH, CANADA INC. | Deprec |

### **Status update – Inactive**

The following MP concepts are generating as active but need to be set to **inactivated**. The DIN is still active but these particular presentations have been discontinued. *This will have to be done manually each month*.

|  |  |  |
| --- | --- | --- |
| **mp\_code** | **mp \_formal\_name** | **mp\_status** |
| 77700067 | DIASTAT (diazepam 2.5 mg per 0.5 mL rectal gel tube) BAUSCH HEALTH, CANADA INC. | Inactive |
| 77700071 | DIASTAT (diazepam 20 mg per 4 mL rectal gel tube) BAUSCH HEALTH, CANADA INC. | Inactive |
| 77700069 | DIASTAT (diazepam 7.5 mg per 1.5 mL rectal gel tube) BAUSCH HEALTH, CANADA INC. | Inactive |

### **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** |
| 12412 | 1958836 | ACETAMINOPHEN ORAL SOL FOR CHILDREN USP (acetaminophen 160 mg per 5 mL oral solution) LABORATOIRES TRIANON INC. | Canceled post-market Exp date: 2022.09.30 |
| 33482 | 2226871 | ACETAMINOPHENE - LIQ 160 MG/5 ML (acetaminophen 160 mg per 5 mL oral solution) LABORATOIRE RIVA INC. | Canceled post-market Exp date: 2022.09.30 |
| 79944 | 2313928 | APO-QUETIAPINE (quetiapine (quetiapine fumarate) 100 mg oral tablet) APOTEX INC | Dormant 2022.08.14 |
| 79943 | 2313901 | APO-QUETIAPINE (quetiapine (quetiapine fumarate) 25 mg oral tablet) APOTEX INC | Dormant 2022.07.08 |
| 79946 | 2313944 | APO-QUETIAPINE (quetiapine (quetiapine fumarate) 300 mg oral tablet) APOTEX INC | Dormant 2022.08.28 |
| 88426 | 2399377 | ATORVASTATIN (atorvastatin (atorvastatin calcium) 10 mg oral tablet) PHARMASCIENCE INC | Canceled post-market Expiry 2021.08.31 |
| 88427 | 2399385 | ATORVASTATIN (atorvastatin (atorvastatin calcium) 20 mg oral tablet) PHARMASCIENCE INC | Canceled post-market Expiry 2021.10.31 |
| 88428 | 2399393 | ATORVASTATIN (atorvastatin (atorvastatin calcium) 40 mg oral tablet) PHARMASCIENCE INC | Canceled post-market Expiry 2021.08.31 |
| 88429 | 2399407 | ATORVASTATIN (atorvastatin (atorvastatin calcium) 80 mg oral tablet) PHARMASCIENCE INC | Canceled post-market Expiry 2021.08.31 |
| 93433 | 2448610 | BYDUREON (exenatide 2 mg powder for prolonged-release suspension for injection with diluent solution per pen) ASTRAZENECA CANADA INC | Canceled post-market Exp date: 2022.04.01 |
| 93036 | 2445131 | CLOTRIMADERM-FLUCONAZOLE COMBI-PACK (clotrimazole 1 % cutaneous cream with fluconazole 150 mg oral capsule) TARO PHARMACEUTICALS INC | Canceled post-market Exp date: 2022.06.30 |
| 70149 | 2246023 | COLD & ALLERGY RELIEF WITH MOISTURIZERS (xylometazoline hydrochloride 0.1 % nasal spray) PHARMASCIENCE INC | Canceled post-market Exp date: 2022.09.30 |
| 89248 | 2407779 | COLD & SINUS RELIEF CAPSULES (ibuprofen 200 mg and pseudoephedrine hydrochloride 30 mg oral capsule) APOTEX INC | Dormant 2022.08.24 |
| 89209 | 2407396 | DIMENHYDRINATE CAPSULES (dimenhydrinate 50 mg oral capsule) APOTEX INC | Dormant 2020.01.22 |
| 97071 | 2480174 | DOVE MEN PLUS CARE DERMA PLUS CARE SCALP ANTI-DANDRUFF DEEP CLEAN 2 IN 1 (pyrithione zinc 1 % shampoo) UNILEVER CANADA | Canceled post-market Exp date: 2022.09.22 |
| 98171 | 2491702 | ERFA HYDROQUINONE (hydroquinone 4 % cutaneous gel) ERFA CANADA 2012 INC | Marketed to Approved. Should be market notification to come. |
| 82345 | 2337819 | EXTAVIA (interferon beta-1b 0.3 mg per vial powder for solution for injection with diluent solution syringe) NOVARTIS PHARMACEUTICALS CANADA INC | Canceled post-market Exp date: 2022.09.30 |
| 89297 | 2408244 | JAMP-LOSARTAN HCTZ (hydrochlorothiazide 12.5 mg and losartan potassium 50 mg oral tablet) JAMP PHARMA CORPORATION | Canceled post-market Exp date: 2022.01.31 |
| 89298 | 2408252 | JAMP-LOSARTAN HCTZ (hydrochlorothiazide 25 mg and losartan potassium 100 mg oral tablet) JAMP PHARMA CORPORATION | Canceled post-market Exp date: 2022.01.31 |
| 91368 | 2429241 | JAMP-RIZATRIPTAN IR (rizatriptan (rizatriptan benzoate) 10 mg oral tablet) JAMP PHARMA CORPORATION | Canceled post-market Exp date: 2022.08.31 |
| 89043 | 2405660 | JAMP-SILDENAFIL (sildenafil (sildenafil citrate) 25 mg oral tablet) JAMP PHARMA CORPORATION | Canceled post-market Exp date: 2021.11.30 |
| 89044 | 2405679 | JAMP-SILDENAFIL (sildenafil (sildenafil citrate) 50 mg oral tablet) JAMP PHARMA CORPORATION | Canceled post-market Exp date: 2021.11.30 |
| 95899 | 2470047 | MILRINONE LACTATE INJECTION (milrinone (milrinone lactate) 10 mg per 10 mL solution for injection vial) AURO PHARMA INC | Dormant 2022.06.10 |
| 95899 | 2470047 | MILRINONE LACTATE INJECTION (milrinone (milrinone lactate) 20 mg per 20 mL solution for injection vial) AURO PHARMA INC | Dormant 2022.06.10 |
| 90931 | 2424630 | NIGHTTIME PAIN RELIEF (diphenhydramine hydrochloride 25 mg and ibuprofen (ibuprofen, ibuprofen potassium) 200 mg oral capsule) APOTEX INC | Dormant 2022.08.06 |
| 99149 | 2501880 | NRA-OMEPRAZOLE (omeprazole (omeprazole magnesium) 20 mg gastro-resistant tablet) NORA PHARMA INC | Cancelled post-market Expiry 2022.09.30 |
| 8054 | 717282 | PRIMAXIN 500 (cilastatin (cilastatin sodium) 500 mg per vial and imipenem 500 mg per vial powder for solution for injection) MERCK CANADA INC | Cancelled post-market Expiry 2022.09.30 |
| 93064 | 2445417 | PRIVA-ROSUVASTATIN (rosuvastatin (rosuvastatin calcium) 5 mg oral tablet) PHARMAPAR INC | Cancelled post-market Expiry 2022.09.30 |
| 79757 | 2312050 | PRO-PIOGLITAZONE (pioglitazone (pioglitazone hydrochloride) 15 mg oral tablet) PRO DOC LIMITEE | Cancelled post-market Expiry 2022.09.30 |
| 79758 | 2312069 | PRO-PIOGLITAZONE (pioglitazone (pioglitazone hydrochloride) 30 mg oral tablet) PRO DOC LIMITEE | Cancelled post-market Expiry 2022.09.30 |
| 6067 | 632775 | RITALIN-SR TAB 20MG (methylphenidate hydrochloride 20 mg prolonged-release oral tablet) NOVARTIS PHARMACEUTICALS CANADA INC | Cancelled post-market Expiry 2022.09.30 |
| 18776 | 2006758 | SOFLAX SYRUP 4MG/ML (docusate sodium 20 mg per 5 mL syrup) PHARMASCIENCE INC | Cancelled post-market 2022.09.15 |
| 85755 | 2372436 | STAXYN (vardenafil (vardenafil hydrochloride) 10 mg orodispersible tablet) BAYER INC | Dormant 2022.08.31 |
| 75035 | 2264846 | TRAMACET (acetaminophen 325 mg and tramadol hydrochloride 37.5 mg oral tablet) JANSSEN INC | Cancelled post-market Expiry 2022.09.30 |
| 16103 | 2014165 | UNIPHYL (theophylline 400 mg prolonged-release oral tablet) ELVIUM LIFE SCIENCES | Cancelled post-market Expiry 2022.09.08 |
| 10314 | 745790 | WHITE PETROLEUM JELLY 100% BP (white petrolatum 100 % cutaneous ointment) H.J. SUTTON INDUSTRIES LTD. | Cancelled post-market Expiry 2022.09.30 |

## **Blacklist (on GitHub)**

The following changes have been made to the Blacklist for the AUGUST 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 |
| d278c98e3c6d9cfb1c0d930571e10b1c | elasomeran 0.1 mg per mL dispersion for injection 2.5 mL vial | 101673 | 2527685 | SPIKEVAX (elasomeran 0.1 mg per mL dispersion for injection 2.5 mL vial) MODERNATX, INC. | REMOVE |
| 9013123 | fludarabine phosphate 50 mg per 2 mL solution for injection vial | 92295 | 2438577 | FLUDARABINE PHOSPHATE INJECTION, USP (fludarabine phosphate 50 mg per 2 mL solution for injection vial) ACCORD HEALTHCARE INC | REMOVE |
| 9014195 | levetiracetam 500 mg per 5 mL solution for injection vial | 101643 | 2527375 | LEVETIRACETAM INJECTION USP (levetiracetam 500 mg per 5 mL solution for injection vial) STERIMAX INC | REMOVE |