

Updates to the Alberta Drug Benefit List

Effective July 1, 2021



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Administered by Alberta Blue Cross
on behalf of Alberta Health.

The Drug Benefit List (DBL) is a list of drugs for which coverage may be provided to program participants. The DBL is not intended to be, and must not be used as a diagnostic or prescribing tool. Inclusion of a drug on the DBL does not mean or imply that the drug is fit or effective for any specific purpose. Prescribing professionals must always use their professional judgment and should refer to product monographs and any applicable practice guidelines when prescribing drugs. The product monograph contains information that may be required for the safe and effective use of the product.

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Special Authorization

The following drug product(s) will be considered for coverage by Special Authorization for patients covered under Alberta government-sponsored drug programs.

New Drug Product(s) Available by Special Authorization

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|-------------------------------------|----------------------------|-------------|------------|
| OLUMIANT 2 MG TABLET | BARCITINIB | 00002480018 | LIL |

Additional Brand(s) and/or Strength(s) of Drug Product(s) Available by Special Authorization

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|-------------------------------------|----------------------------|-------------|------------|
| TEVA-FEBUXOSTAT 80 MG TABLET | FEBUXOSTAT | 00002466198 | TEV |

Additional Brand(s) and/or Strength(s) of Drug Product(s) Available by Restricted Benefit/ Special Authorization

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|---|----------------------------|-------------|------------|
| AG-RIZATRIPTAN ODT 10 MG ORAL DISINTEGRATING TABLET | RIZATRIPTAN BENZOATE | 00002492490 | AGP |

Restricted Benefit(s)

Additional Brand(s) and/or Strength(s) of Drug Product(s) Available by Restricted Benefit

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|-------------------------------------|----------------------------|-------------|------------|
| ARIPIRAZOLE 2 MG TABLET | ARIPIRAZOLE | 00002506688 | SNS |
| ARIPIRAZOLE 5 MG TABLET | ARIPIRAZOLE | 00002506718 | SNS |

Drug Product(s) with Changes to Benefit Status

The following drug product(s) previously covered as regular benefits will now be covered as restricted benefits effective July 1, 2021. Existing patients will be required to transition to the biosimilar by Jan 10, 2022 in order to maintain coverage for the molecule through their Alberta government sponsored drug plan.

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|--|----------------------------|-------------|------------|
| LOVENOX (0.3 ML SYRINGE) 30 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002012472 | SAV |
| LOVENOX (0.4 ML SYRINGE) 40 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002236883 | SAV |
| LOVENOX (0.6 ML SYRINGE) 60 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002378426 | SAV |

Drug Product(s) with Changes to Benefit Status, continued

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|--|----------------------------|-------------|------------|
| LOVENOX (0.8 ML SYRINGE) 120 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002242692 | SAV |
| LOVENOX (0.8 ML SYRINGE) 80 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002378434 | SAV |
| LOVENOX (1 ML SYRINGE) 100 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002378442 | SAV |
| LOVENOX 100 MG / ML INJECTION | ENOXAPARIN SODIUM | 00002236564 | SAV |
| LOVENOX HP (1 ML SYRINGE) 150 MG / SYRINGE INJECTION SYRINGE | ENOXAPARIN SODIUM | 00002378469 | SAV |

Added Product(s)

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|--|------------------------------|-------------|------------|
| AG-URSODIOL 250 MG TABLET | URSODIOL | 00002505363 | AGP |
| AG-URSODIOL 500 MG TABLET | URSODIOL | 00002505371 | AGP |
| ARIPIRAZOLE 10 MG TABLET | ARIPIRAZOLE | 00002506726 | SNS |
| ARIPIRAZOLE 15 MG TABLET | ARIPIRAZOLE | 00002506734 | SNS |
| ARIPIRAZOLE 20 MG TABLET | ARIPIRAZOLE | 00002506750 | SNS |
| ARIPIRAZOLE 30 MG TABLET | ARIPIRAZOLE | 00002506785 | SNS |
| INCLUNOX 40 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002507528 | SDZ |
| INCLUNOX 60 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002507536 | SDZ |
| INCLUNOX 80 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002507544 | SDZ |
| INCLUNOX 100 MG /SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002507552 | SDZ |
| INCLUNOX HP 120 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002507560 | SDZ |
| INCLUNOX HP 150 MG / SYRINGE INJECTION | ENOXAPARIN SODIUM | 00002507579 | SDZ |
| JAMP PILOCARPINE 5 MG TABLET | PILOCARPINE HYDROCHLORIDE | 00002509571 | JPC |
| JAMP SODIUM POLYSTYRENE SULFONATE ORAL POWDER | SODIUM POLYSTYRENE SULFONATE | 00002497557 | JPC |
| JAMP-AMLODIPINE 10 MG TABLET | AMLODIPINE BESYLATE | 00002357208 | JPC |
| MAR-FLECAINIDE 50 MG TABLET | FLECAINIDE ACETATE | 00002476177 | MAR |
| MAR-FLECAINIDE 100 MG TABLET | FLECAINIDE ACETATE | 00002476185 | MAR |
| METHOTREXATE (PRESERVED) 25 MG / ML INJECTION BP | METHOTREXATE SODIUM | 00002464365 | AHI |
| ODAN-SODIUM POLYSTYRENE SULFONATE ORAL POWDER | SODIUM POLYSTYRENE SULFONATE | 00002473941 | ODN |
| REDESCA (3 ML VIAL) 100 MG / ML INJECTION | ENOXAPARIN SODIUM | 00002509121 | VLP |

Added Product(s), continued

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|---|----------------------------|-------------|------------|
| REDESCA 100 MG / ML SINGLE DOSE PRE-FILLED SYRINGE | ENOXAPARIN SODIUM | 00002509113 | SHE |
| REDESCA 30 MG / 0.3 ML SINGLE DOSE PRE-FILLED SYRINGE | ENOXAPARIN SODIUM | 00002509075 | SHE |
| REDESCA 40 MG / 0.4 ML SINGLE DOSE PRE-FILLED SYRINGE | ENOXAPARIN SODIUM | 00002509083 | SHE |
| REDESCA 60 MG / 0.6 ML SINGLE DOSE PRE-FILLED SYRINGE | ENOXAPARIN SODIUM | 00002509091 | SHE |
| REDESCA 80 MG / 0.8 ML SINGLE DOSE PRE-FILLED SYRINGE | ENOXAPARIN SODIUM | 00002509105 | SHE |
| REDESCA HP 120 MG / 0.8 ML SINGLE DOSE PRE-FILLED SYRINGE | ENOXAPARIN SODIUM | 00002509148 | SHE |
| REDESCA HP 150 MG / ML SINGLE DOSE PRE-FILLED SYRINGE | ENOXAPARIN SODIUM | 00002509156 | SHE |
| SANDOZ MIRTAZAPINE 15 MG TABLET | MIRTAZAPINE | 00002250594 | SDZ |

Least Cost Alternative (LCA) Price Change(s)

The following established IC Grouping(s) are affected and a revised LCA price has been established. Groupings affected by a price decrease, will be effective August 1, 2021. Please review the online [Interactive Drug Benefit List](#) for further information.

| <u>Generic Description</u> | <u>Strength / Form</u> | <u>New LCA Price</u> |
|------------------------------|----------------------------------|----------------------|
| AZITHROMYCIN | 20 MG / ML ORAL SUSPENSION | 0.5881 |
| AZITHROMYCIN | 40 MG / ML ORAL SUSPENSION | 0.8330 |
| FEBUXOSTAT | 80 MG TABLET | 0.3975 |
| METHOTREXATE SODIUM | (PRESERVED) 25 MG / ML INJECTION | 6.2400 |
| ONDANSETRON HCL DIHYDRATE | 4 MG TABLET | 2.5450 |
| ONDANSETRON HCL DIHYDRATE | 8 MG TABLET | 3.8840 |
| PILOCARPINE HCL | 5 MG TABLET | 0.7321 |
| RIZATRIPTAN BENZOATE | 5 MG ORAL DISINTEGRATING TABLET | 2.8150 |
| RIZATRIPTAN BENZOATE | 10 MG ORAL DISINTEGRATING TABLET | 2.8150 |
| SODIUM POLYSTYRENE SULFONATE | 94.3 MG / G ORAL POWDER | 0.0648 |

Product(s) with a Price Change

The following product(s) had a Price Change. The previous higher price will be recognized until July 31, 2021. For products within an established IC Grouping, the LCA price may apply.

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|-------------------------------------|----------------------------|-------------|------------|
| ACCEL-ONDANSETRON 4 MG TABLET | ONDANSETRON HCL DIHYDRATE | 00002478927 | ACP |

Product(s) with a Price Change, continued

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|--|------------------------------|-------------|------------|
| ACCEL-ONDANSETRON 8 MG TABLET | ONDANSETRON HCL DIHYDRATE | 00002478935 | ACP |
| M-PILOCARPINE 5 MG TABLET | PILOCARPINE HCL | 00002496119 | ACP |
| ACCEL-RIZATRIPTAN ODT 10 MG ORAL DISINTEGRATING TABLET | RIZATRIPTAN BENZOATE | 00002483289 | ACP |
| ACCEL-RIZATRIPTAN ODT 5 MG ORAL DISINTEGRATING TABLET | RIZATRIPTAN BENZOATE | 00002483270 | ACP |
| AURO-AZITHROMYCIN 20 MG / ML ORAL SUSPENSION | AZITHROMYCIN | 00002482363 | AUR |
| AURO-AZITHROMYCIN 40 MG / ML ORAL SUSPENSION | AZITHROMYCIN | 00002482371 | AUR |
| JAMP-FEBUXOSTAT 80 MG TABLET | FEBUXOSTAT | 00002490870 | JPC |
| MAR-FEBUXOSTAT 80 MG TABLET | FEBUXOSTAT | 00002473607 | MAR |
| METHOTREXATE SODIUM (PRESERVED) 25 MG / ML INJECTION | METHOTREXATE SODIUM | 00002182777 | PFI |
| SANDOZ AZITHROMYCIN 20 MG / ML ORAL SUSPENSION | AZITHROMYCIN | 00002332388 | SDZ |
| SANDOZ AZITHROMYCIN 40 MG / ML SUSPENSION | AZITHROMYCIN | 00002332396 | SDZ |
| SOLYSTAT ORAL POWDER | SODIUM POLYSTYRENE SULFONATE | 0000755338 | PPH |

Discontinued Listing(s)

Notification of discontinuation has been received from the manufacturer(s). The Alberta government-sponsored drug programs previously covered the following drug product(s). Effective July 1, 2021, the listed product(s) will no longer be a benefit and where applicable, will not be considered for coverage by Special Authorization. A transition period will be applied and as of August 1, 2021 claims will no longer pay for these product(s).

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|---|--|-------------|------------|
| ANUGESIC-HC RECTAL OINTMENT | HYDROCORTISONE ACETATE | 00000505781 | MCL |
| APO-QUININE 200 MG CAPSULE | QUININE SULFATE | 00002254514 | APX |
| FAMOTIDINE 20 MG TABLET | FAMOTIDINE | 00002351102 | SNS |
| FAMOTIDINE 40 MG TABLET | FAMOTIDINE | 00002351110 | SNS |
| LOVASTATIN 20 MG TABLET | LOVASTATIN | 00002353229 | SNS |
| LOVASTATIN 40 MG TABLET | LOVASTATIN | 00002353237 | SNS |
| PMS-CARBAMAZEPINE-CR 200 MG SUSTAINED-RELEASE TABLET | CARBAMAZEPINE | 00002231543 | PMS |
| SANDOZ PROCTOMYXIN HC 5 MG / G / 5 MG / G / 10 MG / G / 10 MG / G RECTAL OINTMENT | HYDROCORTISONE/ CINCHOCAINE HCL/ FRAMYCETIN SULFATE/ ESCULIN | 00002242527 | SDZ |
| SANDOZ ZOPICLONE 7.5 MG TABLET | ZOPICLONE | 00002008203 | SDZ |
| TERAZOSIN 1 MG TABLET | TERAZOSIN HCL | 00002350475 | SNS |

Discontinued Listing(s), continued

| <u>Trade Name / Strength / Form</u> | <u>Generic Description</u> | <u>DIN</u> | <u>MFR</u> |
|-------------------------------------|----------------------------|-------------|------------|
| TERAZOSIN 2 MG TABLET | TERAZOSIN HCL | 00002350483 | SNS |

PART 2

Drug Additions

ALBERTA DRUG BENEFIT LIST UPDATE

AMLODIPINE BESYLATE

| 10 MG (BASE) ORAL TABLET | | | | |
|--------------------------|---------------------|-----|----|--------|
| 00002297493 | ACT AMLODIPINE | APH | \$ | 0.1993 |
| 00002331292 | AMLODIPINE | SNS | \$ | 0.1993 |
| 00002385805 | AMLODIPINE | SIV | \$ | 0.1993 |
| 00002429225 | AMLODIPINE | JPC | \$ | 0.1993 |
| 00002419572 | AMLODIPINE BESYLATE | AHI | \$ | 0.1993 |
| 00002273381 | APO-AMLODIPINE | APX | \$ | 0.1993 |
| 00002397080 | AURO-AMLODIPINE | AUR | \$ | 0.1993 |
| 00002357208 | JAMP-AMLODIPINE | JPC | \$ | 0.1993 |
| 00002371723 | MAR-AMLODIPINE | MAR | \$ | 0.1993 |
| 00002362678 | MINT-AMLODIPINE | MPI | \$ | 0.1993 |
| 00002272121 | MYLAN-AMLODIPINE | MYP | \$ | 0.1993 |
| 00002476479 | NRA-AMLODIPINE | NRA | \$ | 0.1993 |
| 00002469049 | PHARMA-AMLODIPINE | PMS | \$ | 0.1993 |
| 00002321866 | RAN-AMLODIPINE | RAN | \$ | 0.1993 |
| 00002284391 | SANDOZ AMLODIPINE | SDZ | \$ | 0.1993 |
| 00002357720 | SEPTA-AMLODIPINE | SEP | \$ | 0.1993 |
| 00000878936 | NORVASC | UJC | \$ | 2.0939 |

ARIPIRAZOLE

| 2 MG ORAL TABLET | | | | |
|------------------|--------------------|-----|----|--------|
| 00002471086 | APO-ARIPIRAZOLE | APX | \$ | 0.8092 |
| 00002506688 | ARIPIRAZOLE | SNS | \$ | 0.8092 |
| 00002460025 | AURO-ARIPIRAZOLE | AUR | \$ | 0.8092 |
| 00002483556 | MINT-ARIPIRAZOLE | MPI | \$ | 0.8092 |
| 00002466635 | PMS-ARIPIRAZOLE | PMS | \$ | 0.8092 |
| 00002473658 | SANDOZ ARIPIRAZOLE | SDZ | \$ | 0.8092 |
| 00002322374 | ABILIFY | OTS | \$ | 3.1618 |

ALBERTA HEALTH RESTRICTED BENEFIT

This Drug Product is a benefit for patients 13 to 17 years of age inclusive.

| 5 MG ORAL TABLET | | | | |
|------------------|--------------------|-----|----|--------|
| 00002471094 | APO-ARIPIRAZOLE | APX | \$ | 0.9046 |
| 00002506718 | ARIPIRAZOLE | SNS | \$ | 0.9046 |
| 00002460033 | AURO-ARIPIRAZOLE | AUR | \$ | 0.9046 |
| 00002483564 | MINT-ARIPIRAZOLE | MPI | \$ | 0.9046 |
| 00002466643 | PMS-ARIPIRAZOLE | PMS | \$ | 0.9046 |
| 00002473666 | SANDOZ ARIPIRAZOLE | SDZ | \$ | 0.9046 |
| 00002322382 | ABILIFY | OTS | \$ | 3.5591 |

ALBERTA HEALTH RESTRICTED BENEFIT

This Drug Product is a benefit for patients 13 to 17 years of age inclusive.

| 10 MG ORAL TABLET | | | | |
|-------------------|--------------------|-----|----|--------|
| 00002471108 | APO-ARIPIRAZOLE | APX | \$ | 1.0754 |
| 00002506726 | ARIPIRAZOLE | SNS | \$ | 1.0754 |
| 00002460041 | AURO-ARIPIRAZOLE | AUR | \$ | 1.0754 |
| 00002483572 | MINT-ARIPIRAZOLE | MPI | \$ | 1.0754 |
| 00002466651 | PMS-ARIPIRAZOLE | PMS | \$ | 1.0754 |
| 00002473674 | SANDOZ ARIPIRAZOLE | SDZ | \$ | 1.0754 |
| 00002322390 | ABILIFY | OTS | \$ | 4.1016 |

ALBERTA DRUG BENEFIT LIST UPDATE

ARIPIPRAZOLE

15 MG ORAL TABLET

| | | | | |
|-------------|----------------------|-----|----|--------|
| 00002471116 | APO-ARIPIPRAZOLE | APX | \$ | 1.2692 |
| 00002506734 | ARIPIPRAZOLE | SNS | \$ | 1.2692 |
| 00002460068 | AURO-ARIPIPRAZOLE | AUR | \$ | 1.2692 |
| 00002483580 | MINT-ARIPIPRAZOLE | MPI | \$ | 1.2692 |
| 00002466678 | PMS-ARIPIPRAZOLE | PMS | \$ | 1.2692 |
| 00002473682 | SANDOZ ARIPIIPRAZOLE | SDZ | \$ | 1.2692 |
| 00002322404 | ABILIFY | OTS | \$ | 4.1016 |

20 MG ORAL TABLET

| | | | | |
|-------------|----------------------|-----|----|--------|
| 00002471124 | APO-ARIPIPRAZOLE | APX | \$ | 1.0017 |
| 00002506750 | ARIPIPRAZOLE | SNS | \$ | 1.0017 |
| 00002460076 | AURO-ARIPIPRAZOLE | AUR | \$ | 1.0017 |
| 00002483599 | MINT-ARIPIPRAZOLE | MPI | \$ | 1.0017 |
| 00002466686 | PMS-ARIPIPRAZOLE | PMS | \$ | 1.0017 |
| 00002473690 | SANDOZ ARIPIIPRAZOLE | SDZ | \$ | 1.0017 |
| 00002322412 | ABILIFY | OTS | \$ | 4.1016 |

30 MG ORAL TABLET

| | | | | |
|-------------|----------------------|-----|----|--------|
| 00002471132 | APO-ARIPIPRAZOLE | APX | \$ | 1.0017 |
| 00002506785 | ARIPIPRAZOLE | SNS | \$ | 1.0017 |
| 00002460084 | AURO-ARIPIPRAZOLE | AUR | \$ | 1.0017 |
| 00002483602 | MINT-ARIPIPRAZOLE | MPI | \$ | 1.0017 |
| 00002466694 | PMS-ARIPIPRAZOLE | PMS | \$ | 1.0017 |
| 00002473704 | SANDOZ ARIPIIPRAZOLE | SDZ | \$ | 1.0017 |
| 00002322455 | ABILIFY | OTS | \$ | 4.1016 |

AZITHROMYCIN

20 MG / ML ORAL SUSPENSION

| | | | | |
|-------------|---------------------|-----|----|--------|
| 00002482363 | AURO-AZITHROMYCIN | AUR | \$ | 0.5881 |
| 00002332388 | SANDOZ AZITHROMYCIN | SDZ | \$ | 0.5881 |
| 00002223716 | ZITHROMAX | PFI | \$ | 1.1310 |

40 MG / ML ORAL SUSPENSION

| | | | | |
|-------------|---------------------|-----|----|--------|
| 00002482371 | AURO-AZITHROMYCIN | AUR | \$ | 0.8330 |
| 00002332396 | SANDOZ AZITHROMYCIN | SDZ | \$ | 0.8330 |
| 00002223724 | ZITHROMAX | PFI | \$ | 1.6026 |

The DBL is not a prescribing or a diagnostic tool. Prescribers should refer to drug monographs and utilize professional judgment.

PRODUCT IS NOT INTERCHANGEABLE

ALBERTA DRUG BENEFIT LIST UPDATE

ENOXAPARIN SODIUM

100 MG / ML INJECTION

| | | | | | |
|-------------------------------------|-------------|---------------------|-----|----|---------|
| <input checked="" type="checkbox"/> | 00002509121 | REDESCA (3 ML VIAL) | VLP | \$ | 17.7100 |
| <input checked="" type="checkbox"/> | 00002236564 | LOVENOX | SAV | \$ | 22.0567 |

RESTRICTED BENEFIT

"Effective July 1, 2021, existing patients currently using the originator biologic drug, Lovenox, will be required to transition to the biosimilar by Jan 10, 2022 in order to maintain coverage for the molecule through their Alberta government sponsored drug plan. All new patient starts for enoxaparin will be covered for the biosimilar. Lovenox will not be eligible for coverage for new enoxaparin starts."

30 MG / SYR INJECTION SYRINGE

| | | | | | |
|-------------------------------------|-------------|--------------------------|-----|----|--------|
| <input checked="" type="checkbox"/> | 00002509075 | REDESCA (0.3 ML SYRINGE) | VLP | \$ | 5.3500 |
| <input checked="" type="checkbox"/> | 00002012472 | LOVENOX (0.3 ML SYRINGE) | SAV | \$ | 6.6170 |

RESTRICTED BENEFIT

"Effective July 1, 2021, existing patients currently using the originator biologic drug, Lovenox, will be required to transition to the biosimilar by Jan 10, 2022 in order to maintain coverage for the molecule through their Alberta government sponsored drug plan. All new patient starts for enoxaparin will be covered for the biosimilar. Lovenox will not be eligible for coverage for new enoxaparin starts."

40 MG / SYR INJECTION SYRINGE

| | | | | | |
|-------------------------------------|-------------|--------------------------|-----|----|--------|
| <input checked="" type="checkbox"/> | 00002507528 | INCLUNOX | SDZ | \$ | 6.6160 |
| <input checked="" type="checkbox"/> | 00002509083 | REDESCA (0.4 ML SYRINGE) | VLP | \$ | 7.0800 |
| <input checked="" type="checkbox"/> | 00002236883 | LOVENOX (0.4 ML SYRINGE) | SAV | \$ | 8.8220 |

RESTRICTED BENEFIT

"Effective July 1, 2021, existing patients currently using the originator biologic drug, Lovenox, will be required to transition to the biosimilar by Jan 10, 2022 in order to maintain coverage for the molecule through their Alberta government sponsored drug plan. All new patient starts for enoxaparin will be covered for the biosimilar. Lovenox will not be eligible for coverage for new enoxaparin starts."

60 MG / SYR INJECTION SYRINGE

| | | | | | |
|-------------------------------------|-------------|--------------------------|-----|----|---------|
| <input checked="" type="checkbox"/> | 00002507536 | INCLUNOX | SDZ | \$ | 9.9240 |
| <input checked="" type="checkbox"/> | 00002509091 | REDESCA (0.6 ML SYRINGE) | VLP | \$ | 10.6300 |
| <input checked="" type="checkbox"/> | 00002378426 | LOVENOX (0.6 ML SYRINGE) | SAV | \$ | 13.2330 |

RESTRICTED BENEFIT

"Effective July 1, 2021, existing patients currently using the originator biologic drug, Lovenox, will be required to transition to the biosimilar by Jan 10, 2022 in order to maintain coverage for the molecule through their Alberta government sponsored drug plan. All new patient starts for enoxaparin will be covered for the biosimilar. Lovenox will not be eligible for coverage for new enoxaparin starts."

ENOXAPARIN SODIUM**80 MG / SYR INJECTION SYRINGE**

| | | | | | |
|-------------------------------------|-------------|--------------------------|-----|----|---------|
| <input checked="" type="checkbox"/> | 00002507544 | INCLUNOX | SDZ | \$ | 13.2320 |
| <input checked="" type="checkbox"/> | 00002509105 | REDESCA (0.8 ML SYRINGE) | VLP | \$ | 14.1600 |
| <input checked="" type="checkbox"/> | 00002378434 | LOVENOX (0.8 ML SYRINGE) | SAV | \$ | 17.6450 |

RESTRICTED BENEFIT

"Effective July 1, 2021, existing patients currently using the originator biologic drug, Lovenox, will be required to transition to the biosimilar by Jan 10, 2022 in order to maintain coverage for the molecule through their Alberta government sponsored drug plan. All new patient starts for enoxaparin will be covered for the biosimilar. Lovenox will not be eligible for coverage for new enoxaparin starts."

100 MG / SYR INJECTION SYRINGE

| | | | | | |
|-------------------------------------|-------------|------------------------|-----|----|---------|
| <input checked="" type="checkbox"/> | 00002507552 | INCLUNOX | SDZ | \$ | 16.5400 |
| <input checked="" type="checkbox"/> | 00002509113 | REDESCA (1 ML SYRINGE) | VLP | \$ | 17.7100 |
| <input checked="" type="checkbox"/> | 00002378442 | LOVENOX (1 ML SYRINGE) | SAV | \$ | 22.0560 |

RESTRICTED BENEFIT

"Effective July 1, 2021, existing patients currently using the originator biologic drug, Lovenox, will be required to transition to the biosimilar by Jan 10, 2022 in order to maintain coverage for the molecule through their Alberta government sponsored drug plan. All new patient starts for enoxaparin will be covered for the biosimilar. Lovenox will not be eligible for coverage for new enoxaparin starts."

120 MG / SYR INJECTION SYRINGE

| | | | | | |
|-------------------------------------|-------------|-----------------------------|-----|----|---------|
| <input checked="" type="checkbox"/> | 00002507560 | INCLUNOX HP | SDZ | \$ | 19.8480 |
| <input checked="" type="checkbox"/> | 00002509148 | REDESCA HP (0.8 ML SYRINGE) | VLP | \$ | 21.2400 |
| <input checked="" type="checkbox"/> | 00002242692 | LOVENOX (0.8 ML SYRINGE) | SAV | \$ | 26.4670 |

RESTRICTED BENEFIT

"Effective July 1, 2021, existing patients currently using the originator biologic drug, Lovenox, will be required to transition to the biosimilar by Jan 10, 2022 in order to maintain coverage for the molecule through their Alberta government sponsored drug plan. All new patient starts for enoxaparin will be covered for the biosimilar. Lovenox will not be eligible for coverage for new enoxaparin starts."

150 MG / SYR INJECTION SYRINGE

| | | | | | |
|-------------------------------------|-------------|---------------------------|-----|----|---------|
| <input checked="" type="checkbox"/> | 00002507579 | INCLUNOX HP | SDZ | \$ | 24.8100 |
| <input checked="" type="checkbox"/> | 00002509156 | REDESCA HP (1 ML SYRINGE) | VLP | \$ | 26.5500 |
| <input checked="" type="checkbox"/> | 00002378469 | LOVENOX HP (1 ML SYRINGE) | SAV | \$ | 33.0850 |

RESTRICTED BENEFIT

"Effective July 1, 2021, existing patients currently using the originator biologic drug, Lovenox, will be required to transition to the biosimilar by Jan 10, 2022 in order to maintain coverage for the molecule through their Alberta government sponsored drug plan. All new patient starts for enoxaparin will be covered for the biosimilar. Lovenox will not be eligible for coverage for new enoxaparin starts."

ALBERTA DRUG BENEFIT LIST UPDATE

FLECAINIDE ACETATE

50 MG ORAL TABLET

| | | | | |
|-------------|-----------------|-----|----|--------|
| 00002275538 | APO-FLECAINIDE | APX | \$ | 0.1389 |
| 00002459957 | AURO-FLECAINIDE | AUR | \$ | 0.1389 |
| 00002493705 | JAMP FLECAINIDE | JPC | \$ | 0.1389 |
| 00002476177 | MAR-FLECAINIDE | MAR | \$ | 0.1389 |

100 MG ORAL TABLET

| | | | | |
|-------------|-----------------|-----|----|--------|
| 00002275546 | APO-FLECAINIDE | APX | \$ | 0.2779 |
| 00002459965 | AURO-FLECAINIDE | AUR | \$ | 0.2779 |
| 00002493713 | JAMP FLECAINIDE | JPC | \$ | 0.2779 |
| 00002476185 | MAR-FLECAINIDE | MAR | \$ | 0.2779 |

FLUTICASONE PROPIONATE

250 MCG / DOSE INHALATION METERED DOSE AEROSOL

| | | | | |
|-------------|---------------------|-----|----|--------|
| 00002503131 | PMS-FLUTICASONE HFA | PMS | \$ | 0.3752 |
|-------------|---------------------|-----|----|--------|

METHOTREXATE SODIUM

25 MG / ML (BASE) INJECTION

| | | | | |
|-------------|-------------------------------|-----|----|--------|
| 00002464365 | METHOTREXATE (PRESERVED) | AHI | \$ | 6.2440 |
| 00002182777 | METHOTREXATE SOD. (PRESERVED) | PFI | \$ | 6.2440 |

MIRTAZAPINE

15 MG ORAL TABLET

| | | | | |
|-------------|--------------------|-----|----|--------|
| 00002411695 | AURO-MIRTAZAPINE | AUR | \$ | 0.0974 |
| 00002496666 | MIRTAZAPINE | SIV | \$ | 0.0974 |
| 00002256096 | MYLAN-MIRTAZAPINE | MYP | \$ | 0.0974 |
| 00002273942 | PMS-MIRTAZAPINE | PMS | \$ | 0.0974 |
| 00002250594 | SANDOZ MIRTAZAPINE | SDZ | \$ | 0.0974 |

ALBERTA DRUG BENEFIT LIST UPDATE

ONDANSETRON HCL DIHYDRATE

4 MG (BASE) ORAL TABLET

| | | | |
|--------------------|--------------------------|------------|------------------|
| 00002478927 | ACCEL-ONDANSETRON | ACP | \$ 2.5450 |
| 00002458810 | CCP-ONDANSETRON | CEL | \$ 2.6790 |
| 00002297868 | MYLAN-ONDANSETRON | MYP | \$ 2.6790 |
| 00002288184 | APO-ONDANSETRON | APX | \$ 3.2720 |
| 00002296349 | CO ONDANSETRON | APH | \$ 3.2720 |
| 00002313685 | JAMP-ONDANSETRON | JPC | \$ 3.2720 |
| 00002371731 | MAR-ONDANSETRON | MAR | \$ 3.2720 |
| 00002305259 | MINT-ONDANSETRON | MPI | \$ 3.2720 |
| 00002417839 | NAT-ONDANSETRON | NTP | \$ 3.2720 |
| 00002421402 | ONDANSETRON | SNS | \$ 3.2720 |
| 00002258188 | PMS-ONDANSETRON | PMS | \$ 3.2720 |
| 00002274310 | SANDOZ ONDANSETRON | SDZ | \$ 3.2720 |
| 00002213567 | ZOFRAN | NOV | \$ 14.9180 |

8 MG (BASE) ORAL TABLET

| | | | |
|--------------------|--------------------------|------------|------------------|
| 00002478935 | ACCEL-ONDANSETRON | ACP | \$ 3.8840 |
| 00002458802 | CCP-ONDANSETRON | CEL | \$ 4.0880 |
| 00002297876 | MYLAN-ONDANSETRON | MYP | \$ 4.0880 |
| 00002288192 | APO-ONDANSETRON | APX | \$ 4.9930 |
| 00002296357 | CO ONDANSETRON | APH | \$ 4.9930 |
| 00002313693 | JAMP-ONDANSETRON | JPC | \$ 4.9930 |
| 00002371758 | MAR-ONDANSETRON | MAR | \$ 4.9930 |
| 00002305267 | MINT-ONDANSETRON | MPI | \$ 4.9930 |
| 00002417847 | NAT-ONDANSETRON | NTP | \$ 4.9930 |
| 00002421410 | ONDANSETRON | SNS | \$ 4.9930 |
| 00002258196 | PMS-ONDANSETRON | PMS | \$ 4.9930 |
| 00002274329 | SANDOZ ONDANSETRON | SDZ | \$ 4.9930 |
| 00002213575 | ZOFRAN | NOV | \$ 22.7660 |

PILOCARPINE HCL

5 MG ORAL TABLET

| | | | |
|--------------------|-------------------------|------------|------------------|
| 00002509571 | JAMP PILOCARPINE | JPC | \$ 0.7321 |
| 00002496119 | M-PILOCARPINE | MTR | \$ 0.7321 |
| 00002216345 | SALAGEN | AMD | \$ 1.1713 |

ALBERTA DRUG BENEFIT LIST UPDATE

RIZATRIPTAN BENZOATE

RESTRICTED BENEFIT - This product is a benefit for patients 18 to 64 years of age inclusive for the treatment of acute migraine attacks in patients where standard therapy has failed. (Refer to Criteria for Special Authorization of Select Drug Products of the List for eligibility in patients 65 years of age and older; and Criteria for Special Authorization of Select Drug Products of the Alberta Human Services Drug Benefit Supplement for eligibility in Alberta Human Services clients.)

| 5 MG (BASE) ORAL DISINTEGRATING TABLET | | | | |
|---|------------------------|-----|----|---------|
| 00002483270 | ACCEL-RIZATRIPTAN ODT | ACP | \$ | 2.8150 |
| 00002458764 | CCP-RIZATRIPTAN ODT | CEL | \$ | 2.9633 |
| 00002379198 | MYLAN-RIZATRIPTAN ODT | MYP | \$ | 2.9633 |
| 00002465086 | JAMP-RIZATRIPTAN ODT | JPC | \$ | 3.7050 |
| 00002462788 | MAR-RIZATRIPTAN ODT | MAR | \$ | 3.7050 |
| 00002436604 | NAT-RIZATRIPTAN ODT | NTP | \$ | 3.7050 |
| 00002393360 | PMS-RIZATRIPTAN RDT | PMS | \$ | 3.7050 |
| 00002442906 | RIZATRIPTAN ODT | SNS | \$ | 3.7050 |
| 00002446111 | RIZATRIPTAN ODT | SIV | \$ | 3.7050 |
| 00002351870 | SANDOZ RIZATRIPTAN ODT | SDZ | \$ | 3.7050 |
| 00002396661 | TEVA-RIZATRIPTAN ODT | TEV | \$ | 3.7050 |
| 00002240518 | MAXALT RPD | MFC | \$ | 16.5163 |
| 10 MG (BASE) ORAL DISINTEGRATING TABLET | | | | |
| 00002483289 | ACCEL-RIZATRIPTAN ODT | ACP | \$ | 2.8150 |
| 00002458772 | CCP-RIZATRIPTAN ODT | CEL | \$ | 2.9633 |
| 00002379201 | MYLAN-RIZATRIPTAN ODT | MYP | \$ | 2.9633 |
| 00002492490 | AG-RIZATRIPTAN ODT | AGP | \$ | 3.7050 |
| 00002465094 | JAMP-RIZATRIPTAN ODT | JPC | \$ | 3.7050 |
| 00002462796 | MAR-RIZATRIPTAN ODT | MAR | \$ | 3.7050 |
| 00002436612 | NAT-RIZATRIPTAN ODT | NTP | \$ | 3.7050 |
| 00002393379 | PMS-RIZATRIPTAN RDT | PMS | \$ | 3.7050 |
| 00002442914 | RIZATRIPTAN ODT | SNS | \$ | 3.7050 |
| 00002446138 | RIZATRIPTAN ODT | SIV | \$ | 3.7050 |
| 00002351889 | SANDOZ RIZATRIPTAN ODT | SDZ | \$ | 3.7050 |
| 00002396688 | TEVA-RIZATRIPTAN ODT | TEV | \$ | 3.7050 |
| 00002240519 | MAXALT RPD | MFC | \$ | 16.5163 |

SODIUM POLYSTYRENE SULFONATE

| ORAL POWDER | | | | |
|--------------------|---------------------------------------|-----|----|--------|
| 00002473941 | ODAN-SODIUM POLYSTYRENE SULFONATE ODN | | \$ | 0.0648 |
| 00000755338 | SOLYSTAT | PPH | \$ | 0.0648 |
| 00002026961 | KAYEXALATE | SAV | \$ | 0.1851 |
| ORAL/RECTAL POWDER | | | | |
| 00002497557 | JAMP SODIUM POLYSTYRENE SULFONATE | JPC | \$ | 0.0648 |

URSODIOL

| 250 MG ORAL TABLET | | | | |
|--------------------|----------------------|-----|----|--------|
| 00002505363 | AG-URSODIOL | AGP | \$ | 0.3818 |
| 00002472392 | JAMP-URSODIOL | JPC | \$ | 0.3818 |
| 00002273497 | PMS-URSODIOL C | PMS | \$ | 0.3818 |
| 00002426900 | URSODIOL TABLETS USP | GLM | \$ | 0.3818 |
| 00002238984 | URSO | AXC | \$ | 1.5469 |
| 500 MG ORAL TABLET | | | | |
| 00002505371 | AG-URSODIOL | AGP | \$ | 0.7242 |
| 00002472406 | JAMP-URSODIOL | JPC | \$ | 0.7242 |
| 00002273500 | PMS-URSODIOL C | PMS | \$ | 0.7242 |
| 00002426919 | URSODIOL TABLETS USP | GLM | \$ | 0.7242 |
| 00002245894 | URSO DS | AXC | \$ | 2.9345 |

PART 3

Special Authorization

BARICITINIB

Rheumatoid Arthritis

"Special authorization coverage may be provided for use in combination with methotrexate for the reduction in signs and symptoms of severely active Rheumatoid Arthritis (RA) in adult patients (18 years of age or older) who are refractory or intolerant to:

- Methotrexate at 20 mg (PO, SC or IM) or greater total weekly dosage (15 mg or greater if patient is 65 years of age or older) for more than 12 weeks. Patients who do not exhibit a clinical response to PO methotrexate or experience gastrointestinal intolerance to PO methotrexate must have a trial of parenteral methotrexate before being accepted as refractory; AND
- Methotrexate with other disease modifying anti-rheumatic agent(s) (minimum 4 month trial) [e.g., methotrexate with hydroxychloroquine or methotrexate with sulfasalazine]; AND
- Leflunomide (minimum 10 week trial at 20 mg daily).

Special authorization coverage of this agent may be provided for use as monotherapy in adult patients for whom methotrexate is contraindicated and/or for those patients who have experienced serious adverse effects.

'Refractory' is defined as lack of effect at the recommended doses and for duration of treatments specified above.

'Intolerant' is defined as demonstrating serious adverse effects or contraindications to treatments as defined in product monographs.

For coverage, this drug must be initiated by a Specialist in Rheumatology ("RA Specialist").

- Initial coverage may be approved for 2 mg once daily for three months.
- Patients will be limited to receiving a one-month supply of baricitinib per prescription at their pharmacy.
- Patients will not be permitted to switch back to baricitinib if they were deemed unresponsive to therapy.

For continued coverage beyond three months, the patient must meet the following criteria:

1) The patient must be assessed by an RA Specialist after the initial three months to determine response.

2) The RA Specialist must confirm in writing that the patient is a 'responder' that meets the following criteria:

- ACR20 OR an improvement of 1.2 units in the DAS28 score [reported to one (1) decimal place]; AND
- An improvement of 0.22 in HAQ score [reported to two (2) decimal places].

It should be noted that the initial score for the DAS28 or HAQ score on record will be rounded to the correct number of decimal places as indicated above.

Following this assessment, continued coverage may be approved for 2 mg once daily for a period of 12 months. Ongoing coverage may be considered only if the following criteria are met at the end of each 12-month period:

1) The patient has been assessed by an RA Specialist to determine response;

2) The RA Specialist must confirm in writing that the patient has maintained a response to therapy as indicated by:

- confirmation of maintenance of ACR20, or
- maintenance of a minimum improvement of 1.2 units in DAS28 score [reported to one (1) decimal place] from baseline.

3) A current HAQ score [reported to two (2) decimal places] must be included with all renewal requests.

It should be noted that the initial score for the DAS28 or HAQ score on record will be rounded to the correct number of decimal places as indicated above.

Coverage cannot be provided for baricitinib when intended for use in combination with a biologic agent or other Janus kinase (JAK) inhibitors."

All requests (including renewal requests) for baricitinib for Rheumatoid Arthritis must be completed using the

ALBERTA DRUG BENEFIT LIST UPDATE
CRITERIA FOR SPECIAL AUTHORIZATION OF SELECT DRUG PRODUCTS

BARICITINIB

Abatacept/Adalimumab/Anakinra/Baricitinib/Certolizumab/Etanercept/Golimumab/Infliximab/Sarilumab/Tocilizumab/Tofacitinib for Rheumatoid Arthritis Special Authorization Request Form (ABC 60027).

2 MG ORAL TABLET

| | | | | |
|-------------|----------|-----|----|---------|
| 00002480018 | OLUMIANT | LIL | \$ | 50.8997 |
|-------------|----------|-----|----|---------|

FEBUXOSTAT

"For the treatment of symptomatic gout in patients with a documented hypersensitivity to allopurinol.

Special authorization may be granted for 6 months."

Please note: Hypersensitivity to allopurinol is a rare condition that is characterized by a major skin manifestation, fever, multi-organ involvement, lymphadenopathy and hematological abnormalities (eosinophilia, atypical lymphocytes). Intolerance or lack of response to allopurinol will not be covered by this criteria.

All requests for febuxostat must be completed using the Febuxostat Special Authorization Request Form (ABC 60037).

The following product(s) are eligible for auto-renewal.

80 MG ORAL TABLET

| | | | | |
|-------------|-----------------|-----|----|--------|
| 00002490870 | JAMP-FEBUXOSTAT | JPC | \$ | 0.3975 |
| 00002473607 | MAR-FEBUXOSTAT | MAR | \$ | 0.3975 |
| 00002466198 | TEVA-FEBUXOSTAT | TEV | \$ | 0.3975 |
| 00002357380 | ULORIC | TAK | \$ | 1.5900 |

**ALBERTA DRUG BENEFIT LIST UPDATE
CRITERIA FOR SPECIAL AUTHORIZATION OF SELECT DRUG PRODUCTS**

RIZATRIPTAN BENZOATE

(Refer to 28:32.28 of the Alberta Drug Benefit List for coverage of patients 18 to 64 years of age inclusive.)

"For the treatment of acute migraine attacks in patients 65 years of age and older where other standard therapy has failed."

"For the treatment of acute migraine attacks in patients 65 years of age and older who have been using rizatriptan benzoate prior to turning 65."

"Special authorization for both criteria may be granted for 24 months."

In order to comply with the first criteria, information is required regarding previous medications utilized and the patient's response to therapy.

The following product(s) are eligible for auto-renewal.

5 MG (BASE) ORAL DISINTEGRATING TABLET

| | | | | |
|--------------------|------------------------------|------------|-----------|---------------|
| 00002483270 | ACCEL-RIZATRIPTAN ODT | ACP | \$ | 2.8150 |
| 00002458764 | CCP-RIZATRIPTAN ODT | CEL | \$ | 2.9633 |
| 00002379198 | MYLAN-RIZATRIPTAN ODT | MYP | \$ | 2.9633 |
| 00002465086 | JAMP-RIZATRIPTAN ODT | JPC | \$ | 3.7050 |
| 00002462788 | MAR-RIZATRIPTAN ODT | MAR | \$ | 3.7050 |
| 00002436604 | NAT-RIZATRIPTAN ODT | NTP | \$ | 3.7050 |
| 00002393360 | PMS-RIZATRIPTAN RDT | PMS | \$ | 3.7050 |
| 00002442906 | RIZATRIPTAN ODT | SNS | \$ | 3.7050 |
| 00002446111 | RIZATRIPTAN ODT | SIV | \$ | 3.7050 |
| 00002351870 | SANDOZ RIZATRIPTAN ODT | SDZ | \$ | 3.7050 |
| 00002396661 | TEVA-RIZATRIPTAN ODT | TEV | \$ | 3.7050 |
| 00002240518 | MAXALT RPD | MFC | \$ | 16.5163 |

10 MG (BASE) ORAL DISINTEGRATING TABLET

| | | | | |
|--------------------|------------------------------|------------|-----------|---------------|
| 00002483289 | ACCEL-RIZATRIPTAN ODT | ACP | \$ | 2.8150 |
| 00002458772 | CCP-RIZATRIPTAN ODT | CEL | \$ | 2.9633 |
| 00002379201 | MYLAN-RIZATRIPTAN ODT | MYP | \$ | 2.9633 |
| 00002492490 | AG-RIZATRIPTAN ODT | AGP | \$ | 3.7050 |
| 00002465094 | JAMP-RIZATRIPTAN ODT | JPC | \$ | 3.7050 |
| 00002462796 | MAR-RIZATRIPTAN ODT | MAR | \$ | 3.7050 |
| 00002436612 | NAT-RIZATRIPTAN ODT | NTP | \$ | 3.7050 |
| 00002393379 | PMS-RIZATRIPTAN RDT | PMS | \$ | 3.7050 |
| 00002442914 | RIZATRIPTAN ODT | SNS | \$ | 3.7050 |
| 00002446138 | RIZATRIPTAN ODT | SIV | \$ | 3.7050 |
| 00002351889 | SANDOZ RIZATRIPTAN ODT | SDZ | \$ | 3.7050 |
| 00002396688 | TEVA-RIZATRIPTAN ODT | TEV | \$ | 3.7050 |
| 00002240519 | MAXALT RPD | MFC | \$ | 16.5163 |