

Appendix III – Criteria for Coverage of Exception Status Drugs

Those whose Coverage of exception status drugs will be approved according to the following criteria upon review of a prescriber's written request. Forms for Exception Status Drug request, which may be used to facilitate the approval process can be found at <https://novascotia.ca/dhw/pharmacare/exception-status-drugs.asp>.

As an alternative to sending a written request to the Pharmacare office, certain exception status drugs have been assigned criteria codes. To allow for online payment of these drugs, the criteria code may be provided by the prescriber on the prescription or confirmed by the pharmacist. The use of these codes offers the prescriber and the pharmacist access to immediate coverage for patients who clearly meet the exception status criteria. The criteria codes are indicated within the following exception criteria.

ABATACEPT (*Orencia 125mg/mL Prefilled Syringe and 250mg/vial Injection*)

- For the treatment of severely active rheumatoid arthritis, in combination with methotrexate or other disease-modifying antirheumatic drugs (DMARDs), in adult patients who are refractory or intolerant to:
 - Methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age), or use in combination with another DMARD, for a minimum of 12 weeks;AND
 - Methotrexate in combination with at least two other DMARDs, such as hydroxychloroquine and sulfasalazine, for a minimum of 12 weeks.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- Optimal treatment response to DMARDs may take up to 24 weeks, however coverage of a biologic therapy can be considered if no improvement is seen after 12 weeks of triple DMARD use.
- If patient factors (e.g. intolerance) prevent the use of triple DMARD therapy, these must be described and dual therapy with DMARDs must be tried.
- 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. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Combined use of more than one biologic DMARD will not be reimbursed.
- Initial Approval: 6 months.
- Renewal Approval: 1 year. Confirmation of continued response is required.
- **Maximum Dosage Approved:**
 - Abatacept Intravenous infusion: 500mg for patients < 60 kg, 750mg for patients 60-100 kg and 1000mg for patients > 100 kg, given at 0, 2, and 4 weeks then every 4 weeks thereafter. Subcutaneous injection: a single IV loading dose of up to 1,000mg may be given, followed by 125mg subcutaneous injection within a day, then once-weekly 125mg subcutaneous injections.
 - Subcutaneous injection: a single IV loading dose of up to 1,000mg may be given, followed by 125mg subcutaneous injection within a day, then once-weekly 125mg subcutaneous injections.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

***ABEMACICLIB** (*Verzenio 50mg, 100mg and 150mg Tablet*)

- In combination with endocrine therapy (ET) for the adjuvant treatment of adult patients with hormone receptor (HR) positive, human epidermal growth factor receptor 2 (HER2) negative, node-positive early breast cancer at high risk of disease recurrence and a Ki-67 score of at least 20%.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should continue until disease progression, unacceptable toxicity, or completion of 2 years of adjuvant therapy. ET may be continued after abemaciclib is completed.
- Patients are not eligible if they have inflammatory breast cancer, or prior treatment with a cyclin-dependent kinases 4 and 6 (CDK4/6) inhibitor.
- Retreatment with a CDK4/6 inhibitor may be reasonable in the metastatic setting if disease recurrence occurs greater than or equal to 6 months after completion of adjuvant abemaciclib.

ABOBOTULINUMTOXIN-A (*Dysport Therapeutic 300U and 500U Vial*)

- For the treatment of cervical dystonia (spasmodic torticollis) in adults.
- For the treatment of upper and lower limb focal spasticity in adults.
- For the treatment of lower limb spasticity in pediatric patients 2 years of age and older.

ABROCITINIB (*Cibinqo 50mg, 100mg and 200mg Tablet*)

- For the treatment of moderate to severe atopic dermatitis (AD) in patients 12 years of age and older who meet all the following criteria:
 - Patients must have had an adequate trial (with a documented refractory disease, including the relief of pruritis), or were intolerant (with documented intolerance), or are ineligible for each of the following therapies:
 - Maximally tolerated medical topical therapies for AD combined with phototherapy (where available), and
 - Maximally tolerated medical topical therapies for AD combined with at least 1 of the 4 systemic immunomodulators (methotrexate, cyclosporine, mycophenolate mofetil, or azathioprine)
 - Baseline Physician Global Assessment score of 3 or greater and Eczema Area and Severity Index (EASI) of 7.1 or greater.

Renewal Criteria:

- Requests for renewal must provide proof of beneficial clinical effect defined as a 75% or greater improvement from baseline in the EASI score (EASI-75) six months after treatment initiation.
- Proof of maintenance of EASI-75 response from baseline must be provided for subsequent authorizations.

Clinical Note:

- Not to be used in combination with phototherapy or any immunomodulatory agents (including biologics or other janus kinase inhibitor treatment) for moderate to severe AD. Treatment should continue until disease progression or unacceptable toxicity.

Claim Notes:

- The patient must be under the care of a dermatologist, allergist, clinical immunologist, or pediatrician who has expertise in the management of moderate to severe AD.
- Approvals will be for a maximum of 200 mg once daily.

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- Initial approval period: 6 months.
- Renewal approval period: 1 year.

***ACALABRUTINIB** (*Calquence 100mg Capsule & Tablet*)

PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)/SMALL LYMPHOCYTIC LYMPHOMA (SLL)

- As a single agent treatment option for adult patients with previously untreated chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) for whom a fludarabine-based regimen is considered inappropriate due to high risk of relapse or refractory disease based on prognostic biomarkers.

Clinical Notes:

- High risk for relapse or refractory disease includes 17p deletion, TP53 mutation, 11q deletion and unmutated IGHV.
- Patients should have a good performance status.
- Treatment should be continued until disease progression or unacceptable toxicity.

Claim Notes:

- Requests will not be considered for patients who experience disease progression on a Bruton's tyrosine kinase (BTK) inhibitor or idelalisib.
- Requests will be considered for patients who are not suitable candidates for intravenous therapy.
- Venetoclax with or without rituximab is funded as a subsequent line of therapy in patients who have experienced disease progression during first-line acalabrutinib treatment, provided all other funding eligibility criteria are met.

RELAPSED/REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA (CLL)/ SMALL LYMPHOCYTIC LYMPHOMA (SLL)

- As a single agent treatment option for adult patients with relapsed or refractory chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) who have received at least one prior therapy.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be continued until disease progression or unacceptable toxicity.

Claim Note:

- Requests will not be considered for patients who experience disease progression on a Bruton's tyrosine kinase (BTK) inhibitor or idelalisib.

ACLIDINIUM BROMIDE (*Tudorza Genuair 400mcg powder for Inhalation*)

- See [Long-Acting Muscarinic Antagonists](#)

ACLIDINIUM/FORMOTEROL (*Duaklir Genuair 12µg/400µg metered dose for Inhalation*)

- See [Long-Acting Beta₂-Agonists/Long-Acting Muscarinic Antagonists](#)

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ADALIMUMAB (*Abrilada; Amgevita; Hulio; Hyrimoz; Idacio; Simlandi; Yuflyma*)
(Please see formulary for specific strength/route(s) covered for each product)

ANKYLOSING SPONDYLITIS

- For the treatment of patients with moderate to severe ankylosing spondylitis (Bath AS Disease Activity Index (BASDAI) score ≥ 4 on 10 point scale) who:
 - Have axial symptoms¹ and who have failed to respond to the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each, or in whom NSAIDs are contraindicated;
 - OR
 - Have peripheral symptoms and who have failed to respond to, or have contraindications to, the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each and have had an inadequate response to an optimal dose or maximal tolerated dose of a DMARD.
- Must be prescribed by a rheumatologist or prescriber with a specialty in rheumatology.
- Requests for renewal must include information showing the beneficial effects of the treatment, specifically:
 - A decrease of at least 2 points on the BASDAI scale, compared with the pre-treatment score; OR
 - Patient and expert opinion of an adequate clinical response as indicated by a significant functional improvement (measured by outcomes such as HAQ or "ability to return to work").

Claim Notes:

- Initial period 6 months.
 - Maximum dose of 40mg every two weeks.
 - Concurrent use of biologics not approved.
1. Patients with recurrent uveitis (2 or more episodes within 12 months) as a complication of axial disease, do not require a trial of 2 NSAIDs.

CROHN'S DISEASE

- For patients with moderate to severely active Crohn's disease and are:
 - Refractory or have contraindications to an adequate course of 5-aminosalicylic acid and corticosteroids and other immunosuppressive therapy;
 - Initial reimbursement is restricted to an induction dose of 160mg followed by 80mg;
 - Clinical response to be assessed twelve weeks after the first induction dose and maintenance therapy approved in responders only at a dose not exceeding 40mg every two weeks.

HIDRADENITIS SUPPURATIVA

- For the treatment of patients with active moderate to severe hidradenitis suppurativa (HS) who have not responded to conventional therapy and who meet all of the following criteria:
 - A total abscess and nodule count of 3 or greater
 - Lesions in at least two distinct anatomic areas, one of which must be Hurley Stage II or III
 - An inadequate response to a 90-day trial of oral antibiotics

Initial renewal criteria:

- Requests for renewal should provide objective evidence of a treatment response, defined as at least a 50% reduction in abscess and inflammatory nodule count with no increase in abscess or draining fistula count relative to baseline at week 12.

Subsequent renewal criteria:

- Requests for renewal should provide objective evidence of the preservation of treatment effect (i.e. the current abscess and inflammatory nodule count and draining fistula count should be compared to the count prior to initiating treatment with adalimumab).

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Claim Notes:

- Must be prescribed by a dermatologist or physician with experience in the treatment of HS.
- Approvals will be for a maximum of 160mg followed by 80mg two weeks later, then 40mg every week beginning four weeks after the initial dose.
- Initial Approval: 12 weeks
- Renewal Approval: 1 year

POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS

- For the treatment of polyarticular juvenile idiopathic arthritis (pJIA) with the following criteria:
 - For patients with moderate or severe pJIA who have had an inadequate response to one or more disease-modifying anti-rheumatic drugs (DMARDs); AND
 - Treatment must be initiated by a rheumatologist who is familiar with the use of DMARDs and/or biologic DMARDs in children.

PSORIASIS

- For patients with severe, debilitating chronic plaque psoriasis who meet all of the following:
 - Body surface area (BSA) involvement of >10% and/or significant involvement of the face, hands, feet or genitals;
 - Failure to, contraindication to or intolerant of methotrexate and cyclosporine;
 - Failure to, intolerant of or unable to access phototherapy;
 - Written request of a dermatologist or prescriber with a specialty in dermatology.
- Continued coverage is dependent on evidence of improvement, specifically:
 - A ≥75% reduction in the Psoriasis Area and Severity Index (PASI) score; or
 - A ≥50% reduction in PASI with a ≥5-point improvement in DLQI (Dermatology Life Quality Index); or
 - Significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals.

Clinical Note:

- Treatment should be discontinued if a response has not been demonstrated after 12 weeks.

Claim Notes:

- Initial approval for a maximum of 16 weeks
- Maximum dosage for ongoing coverage is 40mg every two weeks
- Concurrent use of biologics not approved.

PSORIATIC ARTHRITIS

- For the treatment of patients with predominantly axial psoriatic arthritis who are refractory, intolerant or have contraindications to the sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each.
- For the treatment of patients with predominantly peripheral psoriatic arthritis who are refractory, intolerant or have contraindications to:
 - The sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each; AND
 - Methotrexate (oral or parenteral) at a dose of ≥ 20mg weekly (≥15mg if patient is ≥65 years of age) for a

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minimum of 8 weeks; AND

- Leflunomide for a minimum of 10 weeks or sulfasalazine for a minimum of 3 months.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- 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 to treatments. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial period 3 months.
- Maximum dose of 40mg every two weeks
- Renewal approval: 1 year. Confirmation of continued response required.

RHEUMATOID ARTHRITIS

- For the treatment of severely active rheumatoid arthritis, in combination with methotrexate or other disease-modifying antirheumatic drugs (DMARDs), in adult patients who are refractory or intolerant to:
 - Methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age), or use in combination with another DMARD, for a minimum of 12 weeks;
AND
 - Methotrexate in combination with at least two other DMARDs, such as hydroxychloroquine and sulfasalazine, for a minimum of 12 weeks.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- Optimal treatment response to DMARDs may take up to 24 weeks, however coverage of a biologic therapy can be considered if no improvement is seen after 12 weeks of triple DMARD use.
- If patient factors (e.g. intolerance) prevent the use of triple DMARD therapy, these must be described and dual therapy with DMARDs must be tried.
- 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. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial Approval: 6 months.
- Maximum dose 40mg every two weeks with no dose escalation permitted.
- Renewal Approval: 1 year. Confirmation of continued response is required.

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ULCERATIVE COLITIS

- For the treatment of patients with moderately to severely active ulcerative colitis who have a partial Mayo score > 4, and a rectal bleeding subscore ≥ 2 and are:
 - refractory or intolerant to conventional therapy (i.e. 5-ASA for a minimum of 4 weeks, and prednisone ≥ 40 mg daily for two weeks or IV equivalent for one week); OR
 - corticosteroid dependent (i.e. cannot be tapered from corticosteroids without disease recurrence; or have relapsed within three months of stopping corticosteroids; or require two or more courses of corticosteroids within one year.)
- Renewal requests must include information demonstrating the beneficial effects of the treatment, specifically:
 - a decrease in the partial Mayo score ≥ 2 from baseline, AND
 - a decrease in the rectal bleeding subscore ≥ 1 .

Clinical Notes:

- 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. The nature of intolerance(s) must be clearly documented.
- Patients with severe disease do not require a trial of 5-ASA.

Claim Notes:

- Must be prescribed by a gastroenterologist or physician with a specialty in gastroenterology.
- Concurrent use of biologics not approved.
- Initial Approval: 16 weeks.
- Maximum dose 40mg every two weeks with no dose escalation permitted.
- Renewal Approval: 1 year.

UVEITIS

- For the treatment of patients with non-infectious uveitis who are refractory, intolerant or have contraindications to conventional therapy.

Claim Notes:

- Must be prescribed by, or in consultation with an ophthalmologist.
- Concurrent use of biologics not approved.
- Approvals will be for a maximum of 80 mg followed by 40mg in one week, then 40 mg every two weeks thereafter.
- Initial approval period: 6 months.
- Renewal approval period: 1 year. Confirmation of response is required.

ADEFOVIR DIPVOXIL (*Hepsera 10mg Tablet and generic brands*)

- In combination with lamivudine in patients who:
 - Have developed failure to lamivudine (increase in HBV DNA of ≥ 1 log₁₀iu/mL over the nadir measured on two separate occasions within an interval of at least one month, after the first 3 months of lamivudine therapy); AND
 - When failure to lamivudine is not due to poor adherence to therapy.
- Coverage approved for 1 year.

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*AFATINIB DIMALEATE (*Giotrif 20mg, 30mg, 40mg Tablet*)

- For first line treatment of patients with EGFR mutation positive advanced or metastatic adenocarcinoma of the lung and with an ECOG performance status 0 or 1.

*ALECTINIB (*Alecensaro 150mg Capsule*)

- For the **first line treatment** of patients with locally advanced or metastatic anaplastic lymphoma kinase (ALK) positive non-small cell lung cancer (NSCLC).
- For the treatment of patients with locally advanced or metastatic anaplastic lymphoma kinase (ALK) positive non-small cell lung cancer (NSCLC) who have disease progression on, or intolerance to crizotinib.

Claim Notes:

- Patients should have a good performance status and treatment should be continued until disease progression or unacceptable toxicity.
- If alectinib is chosen as first-line therapy, ceritinib is not funded as a subsequent line of therapy.
- Alectinib is not funded following two prior ALK inhibitor therapies (e.g. crizotinib followed by ceritinib)
- Claims for Alecensaro 150mg capsule that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00904400

ALEMTUZUMAB (*Lemtrada 12mg/1.2mL (10mg/mL) concentrated solution for IV infusion in single-use vials*)

- For the management of adult patients with relapsing-remitting multiple sclerosis (RRMS), with active disease defined by clinical and imaging features, who have had an inadequate response to interferon beta or other disease-modifying therapies, if the following clinical criteria are met:
 - At least two attacks (first episode or relapse) in the previous two years, with at least one attack in the previous year;
 - At least one relapse while on at least six months of a disease modifying therapy within the last 10 years;
 - An Expanded Disability Status Scale (EDSS) score of five (5) or less;
 - Prescribed by a specialist with experience in the treatment of multiple sclerosis.

Claim Notes:

- A maximum of two years of therapy (i.e. two treatment courses; 8 vials) will be reimbursed.
- Claims for Lemtrada 12mg/1.2mL (10mg/mL concentrated solution for IV infusion in single-use vials) that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904161
 - 00904162
 - 00904163
 - 00904164
 - 00904165
 - 00904166
 - 00904167
 - 00904196
 - 00904197

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ALGLUCOSIDASE ALFA (*Myozyme 50mg Powder for Injection*)

- For the treatment of infantile onset Pompe disease in patients who have had the onset of symptoms and confirmed cardiomyopathy before the age of 12 months; AND
- Participation in the long-term evaluation of the efficacy of treatment by periodic medical assessment as outlined in the monitoring of therapy guidelines.
- Initial approval is for 6 months. Continued coverage will be based on evaluation of the efficacy of treatment by regular medical assessment as outlined in the monitoring and discontinuation of therapy guidelines (available from the Pharmacare Office upon request).

Claim Note:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904187
 - 00904188
 - 00904205

ALIROCUMAB (*Praluent 75 mg/mL and 150 mg/mL Prefilled Pen*)

- For the treatment of heterozygous familial hypercholesterolemia (HeFH) in adult patients who require additional lowering of low-density lipoprotein cholesterol (LDL-C) if the following criteria are met:
 - Definite or probable diagnosis of HeFH using the Simon Broome or Dutch Lipid Network criteria or genetic testing; and
 - Patient is unable to reach LDL-C target (less than 2.0 mmol/L or at least a 50% reduction in LDL-C from untreated baseline) despite confirmed adherence to at least 3 months of continuous treatment with:
 - high-dose statin (e.g., atorvastatin 80 mg, rosuvastatin 40 mg) in combination with ezetimibe; or
 - ezetimibe alone if high dose statin is not possible due to rhabdomyolysis, contraindication or intolerance
- **Initial renewal criteria:**
 - A reduction in LDL-C of at least 40% from baseline or has reached a target LDL-C less than 2.0 mmol/L.
- **Subsequent renewal criteria:**
 - The patient continues to maintain a reduction in LDL-C of at least 40% from baseline or has reached a target LDL-C less than 2.0 mmol/L.

Clinical Notes:

- LDL-C levels must be provided.
- Intolerance to high dose statin will be considered if patient has developed documented, myopathy or abnormal biomarkers (i.e. creatinine kinase greater than 5 times the upper limit of normal) after trial of at least two statins and
 - for each statin, dose reduction was attempted rather than statin discontinuation, and intolerance was reversible upon statin discontinuation, but reoccurred with statin re-challenge where clinically appropriate; and
 - at least one statin was initiated at the lowest daily starting dose; and
 - other known causes of intolerance or abnormal biomarkers have been ruled out.
- For patients who cannot take a statin due to an intolerance or contraindication, details must be provided (ie.

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confirmed rhabdomyolysis, active liver disease, unexplained persistent elevations of serum transaminases exceeding three times the upper limit of normal).

- For patients who cannot take ezetimibe due to an intolerance or contraindication, details must be provided.

Claim Notes:

- Maximum dose approved: 300mg every 4 weeks
- Initial approval: 6 months
- Renewal approval: 1 year

ALLERGEN IMMUNOTHERAPY (*Allergy Serum, Pollinex-R Injection*)

- For immunotherapy with specific, standardized allergenic material, administered in high-dose schedules for carefully selected patients with a diagnosis of:
 - IgE mediated anaphylactic reactions to insect stings; or
 - Severe, seasonal (lasting two or more years) or perennial IgE dependent allergic rhinoconjunctivitis when optimal drug therapy and allergen avoidance have not been sufficiently effective in controlling symptoms; or
 - IgE mediated allergic asthma, specifically where there is a clear temporal association between exposure and signs and symptoms of asthma and when optimal drug therapy and avoidance measures have not been sufficiently effective in controlling symptoms.

Note:

- The allergy serum must be dispensed from a pharmacy on prescription from a prescriber. Initial authorization is for two years, and can be continued for up to five years if improvement is noted.

ALMOTRIPTAN (*Axert 6.25mg, 12.5mg Tablet and generic brands*)

- See [Selective 5HT₁ - Receptor Agonists](#)

ALTEPLASE (*Cathflo 2mg Vial*)

- For the treatment of home hemodialysis central venous catheter occlusion.

Clinical Note:

- Not intended for regularly scheduled use.

AMBRISENTAN (*Volibris 5mg, 10mg Tablet and generic brands*)

- For the treatment of patients with at least Class III pulmonary arterial hypertension (PAH), either idiopathic or associated with connective tissue disease who have failed therapy with sildenafil or who have contraindications to sildenafil.
- Diagnosis must be confirmed by right heart catheterization.
- Request must be from a PAH specialist.

AMIFAMPRIDINE (*Firdapse 10mg Tablet*)

- For the treatment of patients with Lambert-Eaton myasthenic syndrome (LEMS) who are 18 years of age and older.

Renewal Criteria:

- Patients should be assessed for a response to treatment within 3 months of initiating amifampridine.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- A response to treatment is defined as an improvement of at least 30% on the 3TUG test.

Claims Notes:

- The patient should be under the care of a neurologist with expertise in managing LEMS.
- Initial Approval: 6 months
- Renewal Approval: long term

AMIFAMPRIDINE (*Ruzurgi 10mg Tablet*)

- For the treatment of patients with Lambert-Eaton myasthenic syndrome (LEMS) who are 6 years of age and older.

Renewal Criteria:

- Patients should be assessed for a response to treatment within 3 months of initiating amifampridine.
- A response to treatment is defined as an improvement of at least 30% on the 3TUG test.

Claims Notes:

- The patient should be under the care of a neurologist with expertise in managing LEMS.
- Initial approval: 6 months
- Renewal approval: Long term

ANAGRELIDE (*Agrylin 0.5mg Capsule and generic brands*)

NOVA SCOTIA SENIORS' PHARMACARE PROGRAM

- For the treatment of essential thrombocythemia (ET) in patients who have:
 - Failed hydroxyurea therapy (does not provide sufficient platelet reduction); or
 - Intolerable side effects from hydroxyurea therapy.

COMMUNITY SERVICES PHARMACARE PROGRAMS

- For the treatment of essential thrombocythemia (ET) as an alternative to hydroxyurea.

ANIFROLUMAB (*Saphnelo 150mg/mL IV Injection*)

- For the treatment of adult patients with active, autoantibody positive, systemic lupus erythematosus (SLE), in addition to standard therapy, who meet all the following criteria:
 - Moderate to severe SLE, defined as Systemic Lupus Erythematosus Disease Activity Index 2000 (SLEDAI-2K) score of at least 6; AND
 - Unable to control their disease while using an oral corticosteroid (OCS) dose of at least 10mg/day of prednisone or equivalent.

Exclusion Criteria:

- Severe or unstable neuropsychiatric SLE
- Active severe SLE nephritis

Initial Renewal Criteria:

- OCS dose decreased to ≤ 7.5 mg/day of prednisone or equivalent, or OCS dose decreased by at least 50% from baseline; AND
- Reduction in disease activity measured by:
 - Reducing the SLEDAI-2K score to 5 or less; OR
 - British Isles Lupus Activity Group (BILAG) improvement in organ systems and no new worsening.

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Subsequent Renewal Criteria:

- The initial response achieved after the first 12 months of therapy has been maintained.

Clinical Notes:

- SLEDAI-2K and BILAG scores must be provided.

Claim Notes:

- Approval: 12 months.
- Patient should be under the care of a physician with expertise in the diagnosis and management of SLE.
- Not to be used in combination with other biologic treatments.

APALUTAMIDE (*Erleada 60mg and 240mg Tablet*)*NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (nmCRPC)**

- In combination with androgen deprivation therapy (ADT) for the treatment of patients with non-metastatic castration-resistant prostate cancer (nmCRPC) who have no detectable distant metastases (M0) by either CT, MRI or technetium-99m bone scan and who are at high risk of developing metastases¹.

¹High risk of developing metastases is defined as a prostate-specific antigen (PSA) doubling time of ≤ 10 months during continuous ADT.

Clinical Notes:

- Patients should have a good performance status and no risk factors for seizures.
- Treatment should continue until radiographic disease progression or unacceptable toxicity.
- Castration-resistance must be demonstrated during continuous ADT and is defined as 3 PSA rises at least one week apart, with the last PSA greater than 2 ng/mL.
- Castrate levels of testosterone must be maintained.
- Patients with N1 disease, pelvic lymph nodes less than 2cm in short axis located below the common iliac vessels, are eligible for apalutamide.
- Apalutamide will not be funded for patients who experience disease progression on enzalutamide or darolutamide.

METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (mCSPC)

- In combination with androgen deprivation therapy (ADT) for the treatment of patients with metastatic castration-sensitive prostate cancer (mCSPC).

Clinical Notes:

- Patients should have a good performance status and no risk factors for seizures.
- Treatment should continue until disease progression or unacceptable toxicity.
- Patients should have had no prior ADT in the metastatic setting, or are within 6 months of initiating ADT in the metastatic setting with no disease progression.
- Patients will be eligible if they received ADT in the non-metastatic setting as long as at least a one year interval has passed since completion.
- Apalutamide will not be funded for patients who experience disease progression on enzalutamide or darolutamide.

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APOMORPHINE HYDROCHLORIDE (*Kynmobi 10mg, 15mg, 20mg, 25mg and 30mg Film*)

- For the acute, intermittent treatment of “OFF” episodes in patients with Parkinson’s Disease (PD) who meet the following criteria:
 - Apomorphine sublingual should only be used as adjunctive therapy in patients who are experiencing “OFF” episodes despite receiving optimized PD therapy (levodopa and derivatives and adjunctive therapy such as dopaminergic agonists or MAO-B inhibitors or amantadine derivatives).

Clinical Notes:

- Treatment should be discontinued unless an improvement of at least 3.25 points is achieved in the Movement Disorders Society Unified Parkinson’s Disease Rating Scale Part III (MDS-UPDRS III) score measured within 30 to 60 minutes after a titrated dose is administered.
- This assessment should occur not more than one year after Kynmobi has been titrated to a stable and tolerated dose.

Claims Notes:

- Approvals will be for a maximum of five films per day or 90 mg in total (whichever is reached first).
- Patients should be under the care of a physician with experience in the diagnosis and management of PD.
- Initial approval: 12 months
- Renewal: long term

***APREPITANT** (*Emend 80mg, 125mg Capsule and Tri-Pack Capsule*)

- In combination with a 5-HT3 antiemetic and dexamethasone for the prevention of acute and delayed nausea and vomiting in patients receiving:
 - highly emetogenic chemotherapy, **[Criteria Code 01]** OR
 - moderately emetogenic chemotherapy who have had inadequate symptom control using a 5-HT3 antagonist and dexamethasone in a previous cycle. **[Criteria Code 02]**

Clinical Notes:

- Highly emetogenic chemotherapy (HEC) may include, but is not limited to: cisplatin regimens, anthracycline and cyclophosphamide combination regimens, and regimens containing carmustine, mechlorethamine, streptozocin, dacarbazine and cyclophosphamide $\geq 1500\text{mg/m}^2$.
- Patients who receive carboplatin-based regimens with $\text{AUC} \geq 4$ are also eligible to receive aprepitant in combination with a 5-HT3 antiemetic and dexamethasone for the primary prevention of acute and delayed nausea and vomiting.

ARIPIPRAZOLE (*Abilify Maintena 300mg and 400mg Prolonged Release Injectible Suspension*)

- For the treatment of patients who are:
 - not adherent to an oral antipsychotic, OR
 - currently receiving a long-acting injectable antipsychotic and require an alternative long-acting injectable antipsychotic.

Claim Note:

- Requests will not be considered for the treatment of psychotic symptoms related to dementia.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

ARTIFICIAL TEARS, PRESERVATIVE FREE (*Celluvisc, Refresh, Refresh Plus, Refresh Tears and Tears Naturale Free*)

- For patients with a diagnosis of dry eye requiring frequent daily doses of artificial tears, to prevent sensitivity to preservatives or in patients in whom preservative sensitivity is suspected;
- Written request from an ophthalmologist or optometrist confirming the diagnosis will be required to initiate coverage.

***ASCIMINIB** (*Scemblix 20mg and 40mg Tablet*)

PHILADELPHIA CHROMOSOME-POSITIVE CHRONIC MYELOID LEUKEMIA (PH+ CML)

- For the treatment of adult patients with Philadelphia chromosome-positive chronic myeloid leukemia (Ph+ CML) in the chronic phase who meet the following criteria:
 - Treatment failure on or intolerance to a minimum of two prior tyrosine kinase inhibitor (TKI) therapies.
 - No evidence of a T315I or V299L mutation.

Clinical Notes:

- Patients should have a good performance status.
- Not for use in the acute phase or blast phase.

ASFOTASE ALFA (*Strensiq 18mg/0.45mL, 28mg/0.7mL, 40mg/1mL and 80mg/0.8mL Single Use Vial*)

- For the treatment of patients with perinatal, infantile, or juvenile-onset hypophosphatasia (HPP).

Clinical Note:

- Eligibility for the treatment of HPP is determined by the Canadian HPP Clinical Expert Committee. Please contact the Nova Scotia Pharmacare Programs via fax at 1-888-594-4440 for the request form.

Claim Notes:

- Must be prescribed by a metabolic specialist with expertise in the diagnosis and management of HPP.
- Claims for Strensiq 18mg/0.45mL, 28mg/0.7mL, 40mg/1mL and 80mg/0.8mL Single Use Vials that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - Strensiq 18mg/0.45mL Single Use Vial
 - 00904483
 - 00904484
 - 00904485
 - Strensiq 28mg/0.7mL Single Use Vial
 - 00904486
 - 00904487
 - 00904488
 - 00904489
 - 00904490
 - Strensiq 40mg/1mL Single Use Vial
 - 00904491
 - 00904492

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- 00904493
- Strensiq 80mg/0.8mL Single Use Vial
 - 00904494
 - 00904495
 - 00904496
 - 00904497
 - 00904498
 - 00904499
 - 00904500
 - 00904501
 - 00904502
 - 00904504
 - 00904505

ASENAPINE (*Saphris 5mg and 10mg SL Tablet*)

- For the acute treatment of manic or mixed episodes associated with bipolar I disorder as either:
 - monotherapy, after a trial of lithium or divalproex sodium has failed, and trials of less expensive atypical antipsychotic agents have failed due to intolerance or lack of response;
 - co-therapy with lithium or divalproex sodium, after trials of less expensive atypical antipsychotic agents have failed due to intolerance or lack of response.

ATOGEPANT (*Qulipta 10mg, 30mg and 60mg Tablet*)

- For the treatment of patients with episodic¹ migraine who have experienced an inadequate response, intolerance, or contraindication to at least two oral prophylactic migraine medications of different classes.

Renewal:

- Proof of beneficial clinical effect, defined as a reduction of at least 50% in the average number of migraine days per month at the time of first renewal compared with baseline.
- For subsequent renewals, proof that the initial 50% reduction in the average number of migraine days per month has been maintained.

Clinical Notes:

- Baseline number of headache and migraine days per month must be provided at the time of initial request.
- ¹Episodic migraine: migraine headaches on at least 4 days per month and less than 15 headache days per month for more than 3 months.

Claim Notes:

- Initial approval: 6 months
- Renewal approval: 1 year
- Concurrent use of more than one calcitonin gene-related peptide (CGRP) inhibitor will not be reimbursed.
- Must be prescribed by a physician who has experience in the management of migraine headaches.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

*AXITINIB (*Inlyta 1mg and 5mg Tablet*)

- For the treatment of patients with advanced or metastatic renal cell carcinoma when used as:
 - first-line therapy in combination with pembrolizumab
OR
 - second-line therapy following disease progression on a vascular endothelial growth factor receptor tyrosine kinase inhibitor (VEGFR TKI)
OR
 - third-line therapy following disease progression on first-line nivolumab and ipilimumab combination therapy, and a second-line VEGFR TKI.
- Patients must have a good performance status. Treatment should be discontinued upon disease progression or unacceptable toxicity.

Clinical Notes:

- Sequential use of axitinib and everolimus is not permitted except in the case of intolerability or contraindication.
- Sequential use of axitinib (as a single agent) and cabozantinib is not permitted except in the case of intolerance or contraindication. Note: Cabozantinib is funded for patients following progression on first-line axitinib + pembrolizumab.
- For patients treated with nivolumab + ipilimumab first-line and VEGF TKI (sunitinib or pazopanib) second line, either cabozantinib **or** axitinib may be used as third-line therapy.
- Axitinib or cabozantinib monotherapy is funded for patients following progression on first-line lenvatinib + pembrolizumab.
- Both clear cell and non-clear cell histology are eligible for treatment.

*AZACYTIDINE (*Onureg 200mg and 300mg Tab*)

ACUTE MYELOID LEUKEMIA

- As maintenance therapy for adult patients with acute myeloid leukemia (AML) who meet all of the following criteria:
 - Intermediate or poor risk
 - Complete remission (CR) or complete remission with incomplete blood count recovery (CRi) following induction therapy, with or without consolidation treatment.
 - Not eligible for hematopoietic stem cell transplantation (HSCT)

Clinical Notes:

- Newly diagnosed includes patients with AML secondary to prior myelodysplastic syndrome (MDS) or chronic myelomonocytic leukemia (CMML).
- Last dose of chemotherapy should be within 4 months of starting azacitidine maintenance.
- Treatment should be discontinued upon disease relapse (i.e., appearance of greater than 5% blasts in the bone marrow or peripheral blood), unacceptable toxicity, or if patient becomes eligible for allogeneic bone marrow or stem cell transplant during the treatment period.

Claim Note:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - Onureg 200mg – 00904806

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Onureg 300mg – 00904807
- Onureg 300mg – 00904815

***AZITHROMYCIN** (*Zithromax POS 100mg/5mL, 200mg/5mL and 250mg, 600mg Tablet and generic brands*)

- The treatment of infections requiring a macrolide antibiotic when the patient has a documented intolerance to clarithromycin. **[Criteria Code 02]**
- The treatment of chlamydia trachomatis as a single dose of 1g. **[Criteria Code 05]**
- The treatment and prevention of mycobacterium avium complex (MAC). **[Criteria Code 06]**
- The treatment of infections requiring a macrolide antibiotic when the patient is taking medications that would significantly interact with erythromycin/clarithromycin. **[Criteria Code 07]**

BENRALIZUMAB (*Fasenra 30mg/mL Prefilled Syringe and Autoinjector*)

- For the adjunctive treatment of severe eosinophilic asthma in adult patients who are inadequately controlled with high dose inhaled corticosteroids and one or more additional asthma controller(s) (e.g., long-acting beta-agonist), and meets one of the following criteria:
 - blood eosinophil count of $\geq 0.3 \times 10^9/L$ within the past 12 months and has experienced two or more clinically significant asthma exacerbations in the past 12 months, OR
 - blood eosinophil count of $\geq 0.15 \times 10^9/L$ and is receiving maintenance treatment with oral corticosteroids (OCS).

Initial Discontinuation Criteria:

- Baseline asthma control questionnaire score has not improved at 12 months since the initiation of treatment, OR
- No decrease in the daily maintenance OCS dose in the first 12 months of treatment, OR
- Number of clinically significant asthma exacerbations has increased within the previous 12 months.

Subsequent Discontinuation Criteria:

- Baseline asthma control questionnaire score achieved after the first 12 months of therapy has not been maintained subsequently, OR
- Reduction in the daily maintenance OCS dose achieved after the first 12 months of treatment is not maintained subsequently, OR
- Number of clinically significant asthma exacerbations has increased within the previous 12 months.

Clinical Notes:

1. A baseline and annual assessment of asthma symptom control using a validated asthma control questionnaire must be provided.
2. High-dose inhaled corticosteroids is defined as greater than or equal to 500 mcg of fluticasone propionate or equivalent daily dose.
3. A clinically significant asthma exacerbation is defined as worsening of asthma such that the treating physician elected to administer systemic glucocorticoids for at least 3 days or the patient visited an emergency department or was hospitalized.

Claim Notes:

- Must be prescribed by a respirologist, clinical immunologist, allergist or internist with experience in treating severe eosinophilic asthma.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Combined use of benralizumab with other biologics used to treat asthma will not be reimbursed.
- Approvals will be for a maximum of 30 mg every four weeks for 12 weeks, then every eight weeks thereafter.
- Initial approval period: 1 year.
- Renewal approval period: 1 year.

***BENZYDAMINE HCL (0.15% Oral Rinse)**

- For oncology patients only. **[Criteria Code 01]**

BETAHISTINE (Serc 16mg, 24mg Tablet and generic brands)

- For the symptomatic treatment of recurrent episodes of vertigo associated with Meniere's disease.

BIMEKIZUMAB (Bimzelx 160mg/mL Prefilled Syringe and 160 mg/mL Autoinjector)

- For patients with severe, debilitating chronic plaque psoriasis who meet all of the following:
 - Body surface area (BSA) involvement of >10% and/or significant involvement of the face, hands, feet or genitals;
 - Failure to, contraindication to or intolerant of methotrexate and cyclosporine;
 - Failure to, intolerant of or unable to access phototherapy;
 - Written request of a dermatologist or prescriber with a specialty in dermatology.
- Continued coverage is dependent on evidence of improvement, specifically:
 - A >75% reduction in the Psoriasis Area and Severity Index (PASI) score; or
 - A >50% reduction in PASI with a > 5 point improvement in DLQI (Dermatology Life Quality Index); or
 - Significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals.

Clinical Note:

- Treatment should be discontinued if a response has not been demonstrated after 16 weeks.

Claim Notes:

- Concurrent use of biologics not approved.
- Approvals will be for 320mg by subcutaneous injection at weeks 0, 4, 8, 12, and 16, followed by maintenance dosing of 320mg every 8 weeks. Maintenance dosing every 4 weeks may be considered for patients >120kg.
- Initial approval period: 16 weeks
- Renewal approval period: 1 year

***BINIMETINIB (Mektovi 15mg Capsule)**

METASTATIC MELANOMA

- In combination with encorafenib for the treatment of patients with BRAF V600 mutation-positive unresectable or metastatic melanoma.

Clinical Notes:

- Patients should have a good performance status.
- If brain metastases are present, patients should be asymptomatic or have stable symptoms.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Notes:

- Binimetinib will not be reimbursed in patients who have progressed on BRAF targeted therapy.
- Requests will be considered for patients who received adjuvant BRAF targeted therapy if disease progression occurred at least 6 months following completion of therapy.

BOSENTAN (*Tracleer 62.5mg, 125mg Tablet and generic brands*)

- Written initial request from a pulmonary arterial hypertension (PAH) specialist only.
- Diagnosis of PAH should be confirmed by right heart catheterization.
- **IPAH (functional class III and IV):**
 - For the treatment of patients with World Health Organization (WHO) functional class III and IV idiopathic pulmonary arterial hypertension (IPAH) who do not demonstrate vasoreactivity on testing or who do demonstrate vasoreactivity on testing but fail a trial of calcium channel blockers (CCB) or are intolerant to CCB.
- **PAH secondary to scleroderma, congenital heart disease or HIV (functional class III and IV):**
 - For the treatment of patients with World Health Organization (WHO) functional class III and IV pulmonary arterial hypertension (PAH) associated with scleroderma, congenital heart disease or HIV who do not respond to conventional therapy.

***BOSUTINIB** (*Bosulif 100mg and 500mg Tablet*)

- For the treatment of adult patients with chronic, accelerated, or blast phase Philadelphia chromosome positive (Ph +) chronic myelogenous leukemia (CML) who have resistance or intolerance to prior tyrosine kinase inhibitor (TKI) therapy.

***BRIGATINIB** (*Alunbrig 30mg, 90mg, 180mg Tablet and Initiation Pack*)**LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER**

- For the first line treatment of patients with locally advanced or metastatic anaplastic lymphoma kinase (ALK) positive non-small cell lung cancer (NSCLC).

Clinical Notes:

- Patients should have a good performance status and treatment should be continued until disease progression or unacceptable toxicity.
- Patients are not eligible for subsequent ALK inhibitor therapy following disease progression on brigatinib.
- Patients may be switched to an alternate ALK inhibitor in the case of intolerance without disease progression.

Claim Notes:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - Alunbrig 30mg Tab – 00904758
 - Alunbrig 90mg Tab – 00904759
 - Alunbrig 180mg Tab – 00904760
 - Alunbrig Initiation Pack – 00904761

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

BRIVARACETAM (*Brivlera 10mg, 25mg, 50mg, 75mg, 100mg Tablet*)

- For the adjunctive treatment of refractory partial-onset seizures (POS) in patients who are currently receiving two or more antiepileptic drugs, and who have had an inadequate response or intolerance to at least three other antiepileptic drugs.

Claim Note:

- The patient must be under the care of a physician experienced in the treatment of epilepsy.

BRODALUMAB (*Siliq 210mg/1.5 mL Prefilled Syringe*)

- For patients with severe, debilitating chronic plaque psoriasis who meet all of the following:
 - Body surface area (BSA) involvement of >10% and/or significant involvement of the face, hands, feet or genitals;
 - Failure to, contraindication to or intolerant of methotrexate and cyclosporine;
 - Failure to, intolerant of or unable to access phototherapy;
 - Written request of a dermatologist or prescriber with a specialty in dermatology.
- Continued coverage is dependent on evidence of improvement, specifically:
 - A >75% reduction in the Psoriasis Area and Severity Index (PASI) score; or
 - A >50% reduction in PASI with a >5-point improvement in DLQI (Dermatology Life Quality Index); or
 - Significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals.

Clinical Note:

- Treatment should be discontinued if a response has not been demonstrated after 12 weeks.

Claim Notes:

- Concurrent use of biologics not approved.
- Initial approval for a maximum of 12 weeks. Renewal approval: 1 year.
- Approvals will be for 210mg at week 0, 1, 2, followed by 210mg every two weeks.

***BUDESONIDE** (*Pulmicort Nebuamps 0.125mg/mL, 0.25mg/mL, 0.5mg/mL Suspension and generic brands*)

- See [Wet Nebulization Solutions](#)

SINONASAL IRRIGATION

- For patients who require budesonide for sinonasal irrigation when it is prescribed by, or in consultation with, a specialist (e.g., ENT, allergists, immunologists).

Claim Notes:

- Initial Approval: 1 year.
- Renewal Approval: Long term

BUDESONIDE/GLYCOPYRRONIUM/ FORMOTEROL FUMERATE DIHYDRATE (*Breztri Aerosphere 182mcg/8.2mcg/5.8mcg Inhaler*)

- For the treatment of chronic obstructive pulmonary disease (COPD), as defined by spirometry, in patients who experience inadequate control while being treated with a long-acting beta-2 agonist/long-acting muscarinic

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

antagonist (LABA/LAMA).

Clinical Notes:

- COPD is defined by spirometry as a post-bronchodilator FEV1/FVC ratio of less than 0.70. Spirometry reports from any point in time will be accepted. If spirometry cannot be obtained, reasons must be clearly explained and other evidence of COPD severity provided (i.e. MRC Dyspnea Scale Score grade).
- Inadequate control while being treated with a LABA/LAMA for at least two months is defined as persistent symptoms or experiencing two or more exacerbations of COPD in the previous year requiring treatment with antibiotics and/or systemic corticosteroids or at least one exacerbation of COPD requiring hospitalization.
- Patients should not be started on a LABA, LAMA and an inhaled corticosteroid (triple inhaled therapy) as initial therapy.

BUPROPION (*Zyban 150mg Tablet*)

- See [Smoking Cessation Therapies](#)

BUROSUMAB (*Crysvita 10mg/mL, 20mg/mL and 30mg/mL Vial*)

Initiation Criteria:

- For the treatment of patients with X-linked hypophosphatemia (XLH) who meet all the following criteria:
 - Initiated in a pediatric patient who is at least one year of age and in whom epiphyseal closure has not yet occurred
 - Fasting hypophosphatemia
 - Normal renal function (defined as a serum creatinine below the age-adjusted upper limit of normal)
 - Radiographic evidence of rickets with a rickets severity score (RSS) of two or greater
 - Confirmed phosphate-regulating endopeptidase homolog, X-linked (PHEX) gene variant in either the patient or in a directly related family member with appropriate X-linked inheritance

Discontinuation Criteria:

- In pediatric patients under 18 years of age in whom epiphyseal closure has not yet occurred and who met the initiation criteria, treatment should be discontinued if:
 - there is no demonstrated improvement in the 12-month RSS total score from baseline RSS total score; or
 - the patient's RSS total score achieved after the first 12 months of therapy has not been maintained subsequently.
- In adolescent patients who are 13 to 17 years of age in whom epiphyseal closure has occurred and who met the initiation criteria and initiated treatment as a pediatric patient, treatment should be discontinued if any of the following occur:
 - Hyperparathyroidism; or
 - Nephrocalcinosis; or
 - Evidence of fracture or pseudo-fracture based on radiographic assessment.
- In adult patients who met the initiation criteria and initiated treatment as a pediatric patient, treatment should be discontinued if any of the following occur:
 - Hyperparathyroidism; or
 - Nephrocalcinosis; or
 - Evidence of fracture or pseudo-fracture based on radiographic assessment.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Notes:

- Requests will not be considered for treatment-naïve adults.
- Must be prescribed by a physician working in a multidisciplinary team of health care providers who are experienced in the diagnosis and management of XLH.
- Approvals for children (1-17 years of age) will be up to a maximum of 90mg every 2 weeks.
- Approvals for adults (18 years of age and older) will be up to a maximum of 90mg every 4 weeks.
- Approval period: 1 year.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - Crysvida Liq Inj 20mg/mL Vial
 - 00904744
 - 00904745
 - 00904746
 - 00904747
 - 00904748
 - Crysvida Liq Inj 30mg/mL Vial
 - 00904749
 - 00904750
 - 00904751
 - 00904752
 - 00904753

BUTORPHANOL (10mg/mL Nasal Spray)

- For the treatment of migraine, upon the request of a neurologist, prescriber with a specialty in neurology or a specialist in pain management, when conventional forms of therapy are ineffective or inappropriate.

CABOZANTINIB (Cabometyx 20mg, 40mg and 60mg Tablet)*ADVANCED OR METASTATIC RENAL CELL CARCINOMA (RCC)**

- For the treatment of patients with advanced or metastatic renal cell carcinoma (RCC) who have received at least one prior vascular endothelial growth factor receptor (VEGFR) tyrosine kinase inhibitor (TKI) therapy. Treatment may continue until clinically meaningful disease progression or unacceptable toxicity.

Clinical Notes:

- Patients with any histology (clear cell or non-clear cell) and IMDC risk are eligible.
- For patients treated with a VEGFR-TKI (sunitinib or pazopanib) first-line, cabozantinib may be used as either a second or third-line treatment option.
- For patients treated with nivolumab + ipilimumab first-line and VEGFR TKI (sunitinib or pazopanib) second-line, either cabozantinib or axitinib monotherapy may be used as third-line therapy.
- Sequential use of axitinib (as monotherapy) and cabozantinib is not permitted except in the case of intolerance or contraindication. Note: Cabozantinib is funded for patients following progression on first-line axitinib + pembrolizumab.
- Axitinib or cabozantinib monotherapy is funded for patients following progression on first-line lenvatinib + pembrolizumab.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

LOCALLY ADVANCED OR METASTATIC DIFFERENTIATED THYROID CARCINOMA (DTC)

- For the treatment of adult patients with locally advanced or metastatic differentiated thyroid carcinoma (DTC) who have progressed on at least one prior line of vascular endothelial growth factor receptor (VEGFR)-targeted tyrosine kinase inhibitor (TKI) therapy.

Clinical Notes:

- Patients should have a good performance status.
- Patients should be refractory to radioactive iodine therapy (RAI-R) or not eligible for radioactive iodine therapy.
- Treatment should continue until disease progression or unacceptable toxicity.
- Patients will be eligible for funding if intolerant to the prior line of VEGFR-targeted TKI therapy.
- Cabozantinib may be used in the third line setting for RET fusion positive patients after progression on or intolerance to selpercatinib.

UNRESECTABLE HEPATOCELLULAR CARCINOMA (HCC)

- For the treatment of patients with unresectable hepatocellular carcinoma (HCC) in the second line setting who have experienced disease progression on sorafenib or lenvatinib and meet all of the following criteria:
 - Child-Pugh class status of A
 - ECOG performance status of 0 or 1
- Treatment should continue until disease progression or unacceptable toxicity.

Clinical Notes:

- Patients with disease progression on regorafenib are not eligible for reimbursement of cabozantinib.
- Patients who are unable to tolerate regorafenib may be switched to cabozantinib if there is no disease progression and provided all other funding criteria are met.
- Patients with disease progression on atezolizumab in combination with bevacizumab are not eligible for reimbursement of cabozantinib.

CALCIPOTRIOL (*Dovonex 50mcg/g Ointment*)

- For the treatment of psoriasis when conventional therapies have been ineffective or inappropriate.

CALCIPOTRIOL/BETAMETHASONE DIPROPIONATE (*Dovobet 0.5mg/g/50mcg/g Gel and generic brands and Enstilar 50mcg/g/ 0.5mg/g Aer Foam*)

- For the treatment of body and scalp psoriasis after failure of a topical steroid and a vitamin D analogue as single agents.

CANAGLIFLOZIN (*Invokana 100mg and 300mg Tablet*)

- For the treatment of Type II diabetes for patients with:
 - Inadequate glycemic control on metformin and a sulfonylurea; and
 - For whom insulin is not an option.

Note:

- 200mg is not a recognized dose; as such a dose of two 100mg tablets will not be funded.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

CANAKINUMAB (*Ilaris 150mg/1mL Solution for Injection*)

- For the treatment of active systemic juvenile idiopathic arthritis, in patients 2 years of age or older, who have an inadequate response or intolerance to systemic corticosteroids (with or without methotrexate) and tocilizumab.

Clinical Note:

- Intolerance is defined as a serious adverse effect as described in the product monograph. The nature of the intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by, or in consultation with, a rheumatologist, who is familiar with the use of biologic DMARDs in children.
- Combined use of more than one biologic DMARD will not be reimbursed.
- Approvals will be for 4 mg/kg for patients > 9 kg, to a maximum of 300mg, administered every four weeks.
- Initial approval period: 16 weeks.
- Renewal approval period: 1 year. Confirmation of continued response is required.
- Claims that exceed \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00903809

***CEFTOLOZANE/TAZOBACTAM** (*Zerbaxa 1g/0.5g Vial*)

- For the treatment of patients with multidrug-resistant gram-negative infections, specifically caused by extended spectrum beta lactamase (ESBL)-producing Enterobacteriaceae and multidrug-resistant Pseudomonas aeruginosa when alternative agents are not an option.

Claim Note:

- Must be prescribed by, or in consultation with, an infectious disease specialist or medical microbiologist.

***CERITINIB** (*Zykadia 150mg Capsule*)

- For the treatment of patients with locally advanced or metastatic anaplastic lymphoma kinase (ALK) positive non-small cell lung cancer (NSCLC) who experience disease progression on, or intolerance to crizotinib.

Claim Notes:

- Patients should have a good performance status and treatment should be continued until disease progression or unacceptable toxicity
- If alectinib is chosen as first-line therapy, ceritinib is not funded as a subsequent line of therapy.
- Disease progression on any other ALK inhibitor in the second-line setting after crizotinib, precludes the use of ceritinib as a subsequent line of therapy.

CERTOLIZUMAB PEGOL (*Cimzia 200mg/mL SC Injection and 200mg/mL Autoinjector Prefilled Pen*)

ANKYLOSING SPONDYLITIS

- For the treatment of patients with moderate to severe ankylosing spondylitis (Bath AS Disease Activity Index (BASDAI) score ≥ 4 on 10 point scale) who:
 - Have axial symptoms¹ and who have failed to respond to the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each, or in whom NSAIDs are contraindicated;

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

OR

- Have peripheral symptoms and who have failed to respond to, or have contraindications to, the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each and have had an inadequate response to an optimal dose or maximal tolerated dose of a DMARD.
- Must be prescribed by a rheumatologist or prescriber with a specialty in rheumatology.
- Requests for renewal must include information showing the beneficial effects of the treatment, specifically:
 - A decrease of at least 2 points on the BASDAI scale, compared with the pre-treatment score; OR
 - Patient and expert opinion of an adequate clinical response as indicated by a significant functional improvement (measured by outcomes such as HAQ or "ability to return to work").

Claim Notes:

- Initial coverage period 6 months. Loading dose of 400mg at Weeks 0, 2 and 4.
 - Maximum maintenance dose of 200mg every 2 weeks or alternatively, 400mg every 4 weeks.
 - Concurrent use of biologics not approved.
1. Patients with recurrent uveitis (2 or more episodes within 12 months) as a complication of axial disease do not require a trial of 2 NSAIDs.

PSORIATIC ARTHRITIS

- For the treatment of patients with predominantly axial psoriatic arthritis who are refractory, intolerant or have contraindications to the sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each.
- For the treatment of patients with predominantly peripheral psoriatic arthritis who are refractory, intolerant or have contraindications to:
 - The sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each; AND
 - Methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age) for a minimum of 8 weeks; AND
 - Leflunomide for a minimum of 10 weeks or sulfasalazine for a minimum of 3 months.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- 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 to treatments. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial coverage period 3 months, loading dose of 400mg at Weeks 0, 2 and 4.
- Maximum maintenance dose of 200mg every 2 weeks or alternatively, 400mg every 4 weeks
- Renewal approval: 1 year. Confirmation of continued response required.

RHEUMATOID ARTHRITIS

- For the treatment of severely active rheumatoid arthritis, in combination with methotrexate or other disease-

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

modifying antirheumatic drugs (DMARDs), in adult patients who are refractory or intolerant to:

- methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age), or use in combination with another DMARD, for a minimum of 12 weeks;

AND

- methotrexate in combination with at least two other DMARDs, such as hydroxychloroquine and sulfasalazine, for a minimum of 12 weeks.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- Optimal treatment response to DMARDs may take up to 24 weeks, however coverage of a biologic therapy can be considered if no improvement is seen after 12 weeks of triple DMARD use.
- If patient factors (e.g. intolerance) prevent the use of triple DMARD therapy, these must be described and dual therapy with DMARDs must be tried.
- 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. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial Approval: 6 months.
- Maximum dose 400mg at weeks 0, 2 and 4 weeks, then 200mg every 2 weeks (or 400mg every 4 weeks) with no dose escalation permitted.
- Renewal Approval: 1 year. Confirmation of continued response is required.

CETIRIZINE (*Reactine 10mg and 20mg Tablet and generic brands*)

- For chronic urticaria, defined as the presence of hives or lesions for longer than six weeks, which has responded to treatment with cetirizine.

CINACALCET (*Sensipar 30mg, 60mg and 90mg Tablet and generic brands*)

- For the treatment of patients with chronic kidney disease on dialysis with severe secondary hyperparathyroidism who:
 - are not responding to optimal doses of Vitamin D analogues or phosphate binders (calcium or non-calcium based) AND are either not a surgical candidate due to surgical or anesthetic risk OR awaiting kidney transplant;
 - in addition laboratory findings must confirm serum phosphate >1.8 mmol/L, serum calcium ≥ 2.2 mmol/L and iPTH >88 pmol/L on more than one occasion at least 6 weeks apart;
 - ongoing laboratory investigations must include serum calcium, albumin, phosphorous weekly for three weeks and iPTH every 6 weeks.
- Coverage for cinacalcet will be renewed if there is a greater than 30% decrease in iPTH after at least 3 months with escalating dose, indicating the patient is responding.
- Approval period 12 months, provided there has been a greater than 30% decrease in iPTH as stated above.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

CIPROFLOXACIN, OPHTHALMIC (*Ciloxan 0.3% Ophthalmic Solution and generic brands and Ointment*)

- See [Fluoroquinolones, Ophthalmic](#)

***CIPROFLOXACIN, ORAL** (*Cipro 100mg/mL Oral Liquid and 250mg, 500mg, 750mg Tablet and generic brands*)

- See [Fluoroquinolones, Oral](#)

CIPROFLOXACIN & DEXAMETHASONE, OTIC (*Ciprodex Otic Suspension and generic brands*)

- For the treatment of patients with acute otitis media with otorrhea through tympanostomy tubes; or with known or suspected tympanic membrane perforation with otorrhea. **[Criteria Code 01]**
- For the treatment of patients with acute otitis externa in the presence of a tympanostomy tube or with known or suspected perforation of the tympanic membrane. **[Criteria Code 02]**

CLADRIBINE (*Mavenclad 10mg Tablet*)

- For the treatment of adult patients with relapsing-remitting multiple sclerosis (RRMS) who meet all the following criteria:
 - Confirmed diagnosis based on McDonald criteria.
 - Has experienced one or more disabling relapses or new MRI activity in the past year.
 - Ambulatory with or without aid (i.e. has a recent Expanded Disability Status Scale (EDSS) score of less than or equal to 6.5).
 - Refractory or intolerant to at least one disease modifying therapy (e.g., interferon, glatiramer, dimethyl fumarate, teriflunomide, ocrelizumab).

Clinical Notes:

- Treatment should be discontinued for patients with an EDSS score of greater than or equal to 7.
- A relapse is defined as the appearance of new or worsening neurological symptoms in the absence of fever or infection, lasting at least 24 hours yet preceded by stability for at least one month and accompanied by new objective neurological findings observed through evaluation by a neurologist.

Claim Notes:

- Must be prescribed by a neurologist with experience in the treatment of multiple sclerosis.
- Approvals will be for 1.75mg/kg to a maximum of 200mg per treatment year.
- Approval period: 2 years
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904524
 - 00904525
 - 00904526

***COBIMETINIB** (*Cotellic 20mg Tablet*)

- For the treatment of patients with BRAF V600 mutation-positive unresectable or metastatic melanoma when used in combination with vemurafenib.

Renewal Criteria:

- Written confirmation that the patient has responded to treatment and there is no evidence of disease progression.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

- Patients must have a good performance status.
- If brain metastases are present, patients should be asymptomatic or have stable symptoms.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.

Claim Note:

- Cobimetinib will not be reimbursed in patients who have progressed on BRAF and/or MEK inhibitor therapy.

***CODEINE, SUSTAINED RELEASE** (*Codeine Contin 50mg, 100mg, 150mg and 200mg Tablet*)

- For the treatment of mild to moderate chronic pain syndrome, if pain has been controlled by doses less than 200mg q12h.
- Patients may be considered candidates if they are achieving good pain control from immediate-release plain codeine preparations but prefer the convenience of a long-acting preparation, or if they are achieving good pain control from acetaminophen or ASA plus codeine preparations but are limited by the acetaminophen content to no greater than 12 tablets per day.
- Not insured for the treatment of acute pain (e.g., post-operative pain).

***CRIZOTINIB** (*Xalkori 200mg and 250mg Capsule*)**ALK-POSITIVE ADVANCED NON-SMALL CELL LUNG CANCER**

- As a first or second-line therapy for patients with ALK-positive advanced non-small cell lung cancer (NSCLC) with ECOG performance status ≤ 2 .

ROS-1 POSITIVE NON-SMALL CELL LUNG CANCER

- For the first-line treatment of patients with ROS-1 positive non-small cell lung cancer (NSCLC).

Clinical Notes:

- Eligible patients should be previously untreated and have a good performance status.
- Treatment may continue until disease progression or unacceptable toxicity.
- Patients with ROS-1 positive NSCLC who are currently receiving first-line chemotherapy or have been previously treated with chemotherapy or immunotherapy will be eligible for treatment with crizotinib.

***CYANOCOBALAMIN, INJECTION** (*Cyanocobalamin, Vitamin B12 100mcg/mL and 1000mcg/mL Injection*)

- For the treatment of documented cyanocobalamin deficiency, when the oral route is inappropriate or contraindicated. (Criteria applies to all Programs.)

***CYANOCOBALAMIN, ORAL** (*Vitamin B12 500mcg and 1,000mcg Tablet*)

- For the treatment of documented cyanocobalamin deficiency in recipients of the Community Services Pharmacare Program, Family Pharmacare Program and Drug Assistance for Cancer Patients; oral cyanocobalamin is fully insured for Seniors' Pharmacare Program.

***CYANOCOBALAMIN, ORAL IN COMBINATION** (*Vitamin B12 1000mcg SL Tablet with Folic Acid*)

- For the treatment of documented cyanocobalamin deficiency in recipients of the Community Services Pharmacare Program, Family Pharmacare Program and Drug Assistance for Cancer Patients; oral cyanocobalamin is fully insured for Seniors' Pharmacare Program.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

CYCLOSPORINE (*Verkazia 0.1% Ophthalmic Emulsion*)

- For the treatment of pediatric patients between the age of 4 and 18 years of age with severe vernal keratoconjunctivitis (VKC) who meet the following criteria:
 - Grade 3 (severe) or 4 (very severe) on the Bonini scale, or
 - Grade 4 (marked) or 5 (severe) on the modified Oxford scale.

Discontinuation Criteria:

- Treatment should be discontinued if no improvement in signs and symptoms of VKC is observed, or
- Treatment should be discontinued if signs and symptoms of VKC have resolved.

Clinical Notes:

- Documentation of the severity of signs and symptoms of VKC at treatment initiation and renewal must be provided.
- Patients previously treated with cyclosporine 0.1% but who discontinued treatment upon resolution of VKC signs and symptoms are eligible to reinstate treatment if signs and symptoms of severe VKC recur and they meet the initiation criteria.

Claim Notes:

- The patient must be under the care of a physician experienced in the diagnosis and management of VKC.
- Initial approval period: 6 months.
- Renewal approval period: 1 year

CYSTEAMINE (*Cystadrops 3.8mg/mL Oph Sol*)

- For the treatment of corneal cystine crystal deposits (CCCDs) in patients 2 years of age and older with cystinosis.

Clinical Note:

- Diagnosis of cystinosis confirmed by cystinosis (lysosomal cystine transporter) gene mutation or elevated white blood cell cystine levels. Documentation must be provided.

Claim Note:

- Must be prescribed by an ophthalmologist experienced in the treatment of CCCDs.

CYSTEAMINE BITARTRATE (*Procysbi 25mg and 75mg Capsule*)

- For the treatment of infantile nephropathic cystinosis with documented cystinosis (lysosomal cystine transporter) gene mutation.

Claim Notes:

- Must be prescribed by, or in consultation with, a prescriber with experience in the diagnosis and management of cystinosis
- Claims for Procysbi 75mg capsule that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904354
 - 00904355

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

*DABRAFENIB (*Tafinlar 50mg and 75mg Capsule*) AND TRAMETINIB (*Mekinist 0.5mg and 2mg Tablet*)

- Dabrafenib-trametinib combination therapy as a first-line BRAF-mutation targeted treatment for patients with BRAF V600 mutation positive, unresectable or metastatic melanoma and who have an ECOG performance status of 0 or 1. Treatment should continue until disease progression. If brain metastases are present, patients should be asymptomatic or have stable symptoms.
- In the event that a patient is initiated on dabrafenib-trametinib combination therapy and has to discontinue one agent due to toxicity, dabrafenib or trametinib monotherapy as a first-line BRAF-mutation targeted treatment for patients with BRAF V600 mutation positive, unresectable or metastatic melanoma and who have an ECOG performance status of 0 or 1, will be funded, should that be the chosen treatment option. Treatment should continue until disease progression. If brain metastases are present, patients should be asymptomatic or have stable symptoms. For clarity, initiation of treatment with dabrafenib or trametinib monotherapy will not be funded.
- For the adjuvant treatment of patients with stage IIIA (limited to lymph node metastases of > 1 mm) to stage IIID (8th edition of American Joint Committee on Cancer [AJCC] staging system) BRAF-mutated (all BRAF V600 mutations) cutaneous melanoma. Disease must be completely resected including in-transit metastases; however, presence of regional lymph nodes with micrometastases after sentinel lymph node biopsy alone is allowed.

Clinical Notes:

- Patients should have a good performance status.
- Treatment with dabrafenib plus trametinib should continue until disease recurrence, unacceptable toxicity, or up to a maximum of 12 months.
- Patients are eligible to receive 12 months of adjuvant treatment with immunotherapy or BRAF targeted therapy. Patients who are unable to tolerate initial adjuvant therapy, within the first 3 months of treatment, may switch to alternate funded treatment, provided criteria are met.
- Patients with mucosal or ocular melanoma are not eligible for treatment with dabrafenib/trametinib.
- Patients who relapse during, or at any time after adjuvant dabrafenib/trametinib therapy, are eligible for treatment with combination immunotherapy (i.e. nivolumab with ipilimumab) in the metastatic setting. Patients who are not candidates for combination immunotherapy are eligible for single agent nivolumab or pembrolizumab immunotherapy in the metastatic setting.
- Re-treatment with BRAF targeted therapy is funded if the treatment-free interval is ≥ 6 months from the completion of adjuvant BRAF therapy.
- For BRAF-positive patients, BRAF-targeted therapy and immunotherapy (including nivolumab plus ipilimumab combination therapy) may be sequenced in either order upon treatment failure, based on clinician assessment.

DALBAVANCIN HYDROCHLORIDE (*Xydalba 500mg Vial*)

- For the treatment of adult patients with acute bacterial skin and skin structure infections (ABSSSI) who meet all the following criteria:
 - known or suspected methicillin-resistant *Staphylococcus aureus* (MRSA) ABSSSI; AND
 - high risk of nonadherence to outpatient antibiotic treatment or high risk of nonadherence to prolonged hospitalization.

Claim Notes:

- Approvals will be for a maximum 1500mg per treatment course.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

DAPAGLIFLOZIN AND METFORMIN HYDROCHLORIDE (*Xigduo 5mg/850mg and 5mg/1000mg Tablet and generic brands*)

- For the treatment of Type II diabetes for patients:
 - who are already stabilized on therapy with dapagliflozin and metformin to replace the individual components of dapagliflozin and metformin; and
 - for whom insulin is not an option

Claim Note:

- Must have met criteria for dapagliflozin.

***DAPTOMYCIN** (*Cubicin RF 500mg/10mL Single-Use Vial*)

- For the treatment of patients with resistant gram-positive infections, including methicillin-resistant *Staphylococcus aureus* (MRSA) who failed to respond, or have a contraindication or intolerance to vancomycin, or for whom IV vancomycin is not appropriate.

Clinical Note:

- Daptomycin is inhibited by pulmonary surfactant and should not be used to treat respiratory tract infections.

Claim Note:

- Must be prescribed by, or in consultation with, an infectious disease specialist or medical microbiologist.

***DARBEPOETIN** (*Aranesp Syringe Injection*)

- For the treatment of transfusion dependent patients with hematologic malignancies who have a baseline anemia of $\leq 90\text{g/L}$ and whose transfusion requirements are ≥ 2 units of packed red blood cells per month over 3 months.
- Initial approval for 6 months with the documentation of dose, hemoglobin and therapeutic outcome (number of transfusions).
- Subsequent 6-month approvals are dependent on evidence of satisfactory clinical response or reduced treatment requirement to less than 2 units of PRBC monthly.

***DAROLUTAMIDE** (*Nubeqa 300mg Tablet*)

NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (nmCRPC)

- In combination with androgen deprivation therapy (ADT) for the treatment of patients with non-metastatic castration-resistant prostate cancer (nmCRPC) who have no detectable distant metastases (M0) by either CT, MRI or technetium-99m bone scan and who are at high risk of developing metastases¹.

¹High risk of developing metastases is defined as a prostate-specific antigen (PSA) doubling time of ≤ 10 months during continuous ADT.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should continue until radiographic disease progression or unacceptable toxicity.
- Castration-resistance must be demonstrated during continuous ADT and is defined as 3 PSA rises at least one week apart, with the last PSA greater than 2 ng/mL.
- Castrate levels of testosterone must be maintained.
- Patients with N1 disease, pelvic lymph nodes less than 2cm in short axis located below the aortic bifurcation are eligible for darolutamide.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Darolutamide will not be funded for patients who experience disease progression on apalutamide or enzalutamide.

METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (MCSPC)

- In combination with docetaxel and androgen deprivation therapy (ADT) for the treatment of patients with metastatic castration-sensitive prostate cancer (mCSPC).

Clinical Notes:

- Patients should have a good performance status and be eligible for chemotherapy.
- Treatment should continue until disease progression or unacceptable toxicity.
- Patients should have had no prior ADT in the metastatic setting, or are within 6 months of initiating ADT in the metastatic setting with no disease progression.
- Patients will be eligible if they received ADT in the non-metastatic setting as long as at least a one year interval has passed since completion.
- Darolutamide will not be funded for patients who experience disease progression on enzalutamide or apalutamide.

DARIFENACIN (*Enablex 7.5mg, 15mg Tablet and generic brands*)

- See [OAB Medications](#)

***DASATINIB** (*Sprycel 20mg, 50mg, 70mg, 100mg Tablet and generic brands*)

- For the treatment of adult patients with Philadelphia chromosome-positive chronic myeloid leukemia (Ph+ CML) in chronic, accelerated, or blast phase.
- For the treatment of patients with Philadelphia chromosome-positive acute lymphoblastic leukemia (Ph+ ALL).

***DECITABINE AND CEDAZURIDINE** (*Inqovi 35mg/100mg Tablet*)

MYELOYDYSPLASTIC SYNDROME

- For the treatment of patients with myelodysplastic syndromes (MDS), including previously treated and untreated, who meet all of the following criteria:
 - De novo or secondary MDS including all French-American-British subtypes (i.e., refractory anemia, refractory anemia with ringed sideroblasts, refractory anemia with excess blasts, refractory anemia with excess blasts in transformation, and chronic myelomonocytic leukemia)
 - Intermediate-1, intermediate-2, or high-risk MDS, according to the International Prognostic Scoring System
 - Have not experienced disease progression on a hypomethylating agent

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.

DEFERASIROX (*Exjade 125mg, 250mg, 500mg Tablet for Suspension and generic brands*)

- For the treatment of chronic iron overload.

DEFERASIROX (*Jadenu 90mg, 180mg, 360mg Tablet and generic brands*)

- For the treatment of chronic iron overload.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

DEFERIPRONE (*Ferriprox 100mg/mL Solution and 1000mg Tablet*)

- For the treatment of patients with transfusional iron overload due to thalassemia syndromes when current chelation therapy is inadequate.

***DENOSUMAB** (*Prolia 60mg/mL Prefilled Syringe*)

- For the treatment of osteoporosis in postmenopausal women and in men who meet the following criteria:
 - Have a contraindication to oral bisphosphonates; and
 - High risk for fracture, or refractory or intolerant to other available osteoporosis therapies.

Clinical Notes:

- Refractory is defined as a fragility fracture or evidence of a decline in bone mineral density below pre-treatment baseline levels, despite adherence for one year to other available osteoporosis therapies.
- High fracture risk is defined as:
 - Moderate 10-year fracture risk (10% to 20%) as defined by the Canadian Association of Radiologists and Osteoporosis Canada (CAROC) tool or the World Health Organization's Fracture Risk Assessment (FRAX) tool with a prior fragility fracture; or
 - High 10-year fracture risk ($\geq 20\%$) as defined by the CAROC or FRAX tool.

***DENOSUMAB** (*Xgeva 120mg/1.7mL Solution*)

- As a single agent for the prevention of skeletal related events (SREs) for metastatic castrate resistant prostate cancer (CRPC) patients with one or more documented bone metastases and ECOG performance status (PS) 0-2.

DIENOGEST (*Visanne 2mg Tablet and generic brands*)

- For the management of pelvic pain associated with endometriosis in patients for whom one or more less costly hormonal options are either ineffective or cannot be used.

DIMETHYL FUMARATE (*Tecfidera 120mg and 240mg DR Capsule and generic brands*)

- For the treatment of patients with relapsing remitting multiple sclerosis (RRMS) who meet all of the following criteria:
 - requested and followed by a neurologist experienced in the management of RRMS; and
 - recent expanded disability status scale (EDSS) score of 5.5 or less (i.e. patients must be able to ambulate at least 100 metres without assistance).
- **Exclusion:**
 - not funded in combination with other disease modifying therapies;
 - not funded in patients with an EDSS > 5.5 ;
 - not funded in patients < 18 years of age.
- **Renewals:**
 - EDSS score < 5.5 (i.e. patients must be able to ambulate at least 100 metres without assistance). Date and details of the most recent neurological examination and EDSS score must be provided (exam must have occurred within the last 90 days); and
 - patients must be stable or have experienced no more than 1 disabling attack/relapse in the past year.

DIPYRIDAMOLE & ASA (*Aggrenox 200/25mg Capsule and generic brands*)

- For the secondary prevention of ischemic stroke/transient ischemic attack (TIA) in patients who have experienced

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

a recurrent thrombotic event (stroke, symptoms of TIA) while taking ASA.

DUPIUMAB (*Dupixent 200mg/1.14mL Prefilled Syringe/Pen and 300mg/2mL Prefilled Syringe/Pen*)

ATOPIC DERMATITIS

- For the treatment of moderate to severe atopic dermatitis (AD) in patients 12 years of age and older who meet all of the following criteria:
 - Patients must have had an adequate trial (with a documented refractory disease), or were intolerant (with documented intolerance), or are ineligible for each of the following therapies:
 - maximally tolerated medical topical therapies for AD combined with phototherapy (where available), and;
 - maximally tolerated medical topical therapies for AD combined with at least 1 of the 4 systemic immunomodulators (methotrexate, cyclosporine, mycophenolate mofetil, or azathioprine).
 - Baseline Physician Global Assessment score of 3 or greater and Eczema Area and Severity Index (EASI) of 7.1 or greater.

Renewal criteria:

- Requests for renewal must provide proof of beneficial clinical effect defined as a 75% or greater improvement from baseline in the Eczema Area and Severity Index (EASI-75) score six months after treatment initiation.
- Proof of maintenance of EASI-75 response from baseline must be provided for subsequent authorizations.

Clinical Note:

- Not to be used in combination with phototherapy or any immunomodulatory drugs (including biologics) or a Janus kinase inhibitor treatment for moderate-to-severe AD.

Claim Notes:

- The patient must be under the care of a dermatologist, allergist, clinical immunologist, or pediatrician who has expertise in the management of moderate to severe AD.
- Approvals will be for a maximum of 600 mg at week 0, then 300 mg every two weeks thereafter.
- Initial approval period: 6 months.
- Renewal approval period: 1 year.

SEVERE ASTHMA (PEDIATRIC)

- For the adjunctive treatment of severe asthma with a type 2 or eosinophilic phenotype in patients aged 6 to 11 years of age who are inadequately controlled with medium-to high-dose inhaled corticosteroids (ICS) plus one or more additional asthma controller(s) (e.g., long-acting beta-agonist) or high-dose ICS alone and meet the following criteria:
 - blood eosinophil count $\geq 0.15 \times 10^9/L$ within the past 12 months; and
 - uncontrolled asthma with at least one clinically significant asthma exacerbation in the past 12 months.

Initial Discontinuation Criteria:

- Baseline asthma control questionnaire score has not improved at 12 months since initiation of treatment, or
- The number of clinically significant asthma exacerbations has increased within the previous 12 months

Subsequent Discontinuation Criteria:

- Asthma control questionnaire score achieved after the first 12 months of therapy has not been maintained subsequently, or
- The number of clinically significant asthma exacerbations has increased within the previous 12 months.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

- A baseline and annual assessment of asthma symptom control using a validated asthma control questionnaire must be provided.
- Medium dose ICS is defined as between 200 mcg and 400 mcg of fluticasone propionate or equivalent daily dose and high-dose ICS is defined as greater than 400 mcg of fluticasone propionate or equivalent daily dose.
- A significant clinical exacerbation is defined as worsening of asthma such that the treating physician elected to administer systemic glucocorticoids for at least 3 days or the patient visited an emergency department or was hospitalized.

Claim Notes:

- Must be prescribed by a pediatric respirologist or allergist experienced in the treatment of severe asthma.
- Combined use of dupilumab with other biologics used to treat asthma will not be reimbursed.
- Approvals will be for a maximum of 200 mg every two weeks or 300 mg every four weeks.
- Approval period: 1 year.

SEVERE ASTHMA

- For the adjunctive treatment of severe asthma with a type 2 or eosinophilic phenotype or oral corticosteroid (OCS) dependent severe asthma in patients 12 years of age and older who are inadequately controlled with high-dose inhaled corticosteroids (ICS) and one or more additional asthma controller(s) (e.g., long-acting beta-agonist) and meets one of the following criteria:
 - blood eosinophil count $\geq 0.15 \times 10^9/L$ within the past 12 months, or
 - have OCS dependent asthma.

Initial Discontinuation Criteria:

- Baseline asthma control questionnaire score has not improved at 12 months since initiation of treatment, or
- No decrease in the daily maintenance OCS dose in the first 12 months of treatment, or
- Number of clinically significant asthma exacerbations has increased within the previous 12 months.

Subsequent Discontinuation Criteria:

- Asthma control questionnaire score achieved after the first 12 months of therapy has not been maintained subsequently, or
- Reduction in the daily maintenance OCS dose achieved after the first 12 months of treatment is not maintained subsequently, or
- Number of clinically significant asthma exacerbations has increased within the previous 12 months.

Clinical Notes:

- A baseline and annual assessment of asthma symptom control using a validated asthma control questionnaire must be provided.
- A baseline and annual number of clinically significant asthma exacerbations must be provided.
- High-dose ICS is defined as greater than or equal to 500 mcg of fluticasone propionate or equivalent daily dose.
- A significant clinical exacerbation is defined as worsening of asthma such that the treating physician elected to administer systemic glucocorticoids for at least 3 days or the patient visited an emergency department or was hospitalized.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Notes:

- Must be prescribed by a respirologist, clinical immunologist, allergist or internist experienced in the treatment of severe asthma.
- Combined use of dupilumab with other biologics used to treat asthma will not be reimbursed.
- Approvals will be for a maximum of 600 mg at week 0, then 300 mg every two weeks thereafter.
- Approval period: 1 year.

EDARAVONE (*Radicava 30mg/100mL IV Injection and 105mg/5mL Suspension*)

For the treatment of amyotrophic lateral sclerosis (ALS), if the following criteria are met:

Initiation Criteria:

- Patient with a diagnosis of probable ALS or definite ALS; AND
- Patient who meets all of the following:
 - has scores of at least two points on each item of the ALS Functional Rating Scale – Revised (ALSFERS-R)
 - has a forced vital capacity greater than or equal to 80% of predicted
 - has had ALS symptoms for two years or less
 - patient is not currently requiring permanent non-invasive or invasive ventilation.

Renewal Criteria:

- Reimbursement of treatment should be discontinued in patients who meet any one of the following criteria:
 - patient becomes non-ambulatory (ALSFERS-R score \leq 1 for item 8) AND is unable to cut food and feed themselves without assistance, irrespective of whether a gastrostomy is in place (ALSFERS-R score $<$ 1 for item 5a or 5b);
 - OR
 - patient requires permanent non-invasive or invasive ventilation.

Claim Notes:

- Patient must be under the care of a specialist with experience in the diagnosis and management of ALS.
- Claims for Radicava 30mg/100mL IV Injection that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00904538
- Claims for Radicava 105mg/5mL Suspension that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00904996

ELOSULFASE ALFA (*Vimizim 1mg/mL IV Solution*)**MUCOPOLYSACCHARIDOSIS TYPE IVA****Initiation Criteria:**

- For the treatment of mucopolysaccharidosis type IVA (MPS IVA) in patients meeting all the following criteria:
 - Diagnosis is confirmed by enzymatic assay for N-acetylgalactosamine-6-sulfate sulfatase (GALNS) activity in peripheral blood leukocytes or fibroblasts (excluding multiple sulfatase deficiency) AND mutational analysis of GALNS¹; AND
 - Patient is under the care of a specialist with experience in the diagnosis and management of MPS IVA; AND
- The following baseline evaluations prior to initiation of elosulfase alfa must be provided with the request for

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

coverage:

- Detailed medical history documenting surgeries, medical admissions, subspecialty assessments
- Orthopedic evaluation including spinal and cranial MRI, skeletal x-rays, pain symptoms from bone and joints as appropriate to age and clinical disease
- Mobility measure: 6MWT and stair climb (if appropriate for age and disease status)
- Respiratory function testing including sleep study testing (if appropriate for age)
- Age-appropriate quality of life measure (such as HAQ, PODCI, EQ5D5L or SF36)²
- Documentation of mobility aide requirement, such as a walker or cane
- Documentation of requirement for respiratory aides, including ventilation status and changes in respiratory support requirements
- Ophthalmologic and ear, nose and throat (ENT) assessment (if appropriate)
- Urine keratin sulfate (KS) determination: specific KS determination is preferred over total glycosaminoglycans (GAGs)
- Cardiac echocardiogram

¹Note: not all MPS IVA patients will have two known pathogenic alleles identified and parental mutation analysis to establish the phase of mutations should be performed.

²Note that academic goals (e.g. attendance or participation in school) may be considered case-by-case in pediatric patients.

Exclusion Criteria:

- The patient is diagnosed with an additional progressive life limiting condition where treatment would not provide long term benefit (such as cancer or multiple sclerosis).
- The patient has a forced vital capacity (FVC) of less than 0.3 liters and requires continuous ventilator assistance.
- The patient/family is unwilling to comply with the associated monitoring criteria.
- The patient/family is unwilling to attend clinics for assessment and treatment purposes.

Approval duration of initial approval: 1 year

Recommended dose: 2mg/kg IV infusion once a week.

Renewal criteria:

- Patients must demonstrate at least 3 of the 5 following treatment effects for continuation of coverage of treatment with elosulfase alfa:
 - 6 MWT or Stair Climb test stabilized at or improved by at least 5% of baseline measure
 - FVC or forced expiratory volume in one second (FEV-1) stabilized at or improved by at least 5% of baseline measure or remaining within 2 standard deviations of normal for the patient's age
 - Improvement or no change (if minimal effect) in age-appropriate quality of life measure³
 - Reduction of urine KSs of 20%
 - Stability of cardiac ejection fraction reduction (within 5% of baseline)

³Note that academic goals (e.g. attendance or participation in school) may be considered case-by-case in pediatric patients.

Discontinuation criteria:

- Patients will not be eligible for coverage of treatment if they:
 - Fail to meet 3 of the 5 renewal criteria
 - Are unable to tolerate infusions due to infusion related adverse events that cannot be resolved
 - Require permanent invasive ventilation

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Miss more than 6 infusions in a 12-month interval, unless for medically related issues
- Meet any one of the Exclusion Criteria

Approval duration of renewals: 1 year

Recommended dose: 2mg/kg IV infusion once a week.

Claim Note:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904541
 - 00904542
 - 00904543
 - 00904544
 - 00904545

EMPAGLIFLOZIN (*Jardiance 10mg and 25mg Tablet*)

- For the treatment of Type 2 diabetes mellitus for patients with:
 - inadequate glycemic control on metformin and a sulfonylurea; and
 - for whom insulin is not an option

OR
- As an adjunct to diet, exercise, and standard care therapy to reduce the incidence of cardiovascular death in patients with type 2 diabetes mellitus and established cardiovascular disease (details must be provided as per clinical note below) who have:
 - inadequate glycemic control despite an adequate trial of metformin

Clinical Notes:

- Established cardiovascular disease is defined as one of the following (details must be provided):
 - History of myocardial infarction (MI)
 - Multi-vessel coronary artery disease in two or more major coronary arteries (irrespective of revascularization status)
 - Single-vessel coronary artery disease with significant stenosis and either a positive non-invasive stress test or discharged from hospital with a documented diagnosis of unstable angina within 12 months prior to selection.
 - Last episode of unstable angina >2 months prior with confirmed evidence of coronary multi-vessel or single-vessel disease.
 - History of ischemic or hemorrhagic stroke.
 - Occlusive peripheral artery disease.

EMPAGLIFLOZIN/METFORMIN HYDROCHLORIDE (*Synjardy 5mg/500mg, 5mg/850mg, 5mg/1000mg, 12.5mg/500mg, 12.5mg/850mg and 12.5mg/1000mg Tablet*)

- For the treatment of type 2 diabetes mellitus in patients who are already stabilized on therapy with empagliflozin and metformin, to replace the individual components of empagliflozin and metformin. Patients must meet coverage criteria for empagliflozin.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

EMTRICITABINE/TENOFOVIR DISOPROXIL FUMARATE (*Truvada 200mg/ 300mg Tablet and generic brands*)

MEN WHO HAVE SEX WITH MEN (MSM) AND TRANSGENDER WOMEN (TGW)

- For pre-exposure prophylaxis (PrEP), in combination with safer sex practices, to reduce the risk of sexually acquired HIV-1 infection in adults at high risk who report condomless anal sex within the last six months and any of the following:
 - Infectious syphilis or rectal bacterial sexually transmitted infection (STI), particularly if diagnosed in the preceding 12 months;
 - Recurrent use of nonoccupational postexposure prophylaxis (nPEP) (more than once);
 - Ongoing sexual relationship with an HIV-positive partner who is not receiving stable ART and/or does not have an HIV viral load <200 copies/ mL. (i.e. not on ART or >200 copies/mL); or
 - High-incidence risk index (HIRI)-MSM risk score ≥ 11 . Please refer to the [BC-CfE PrEP guidelines](#) or the [Canadian PrEP Guidelines](#) which include details about how to calculate the HIRI-MSM risk score.

HETEROSEXUAL EXPOSURE

- For pre-exposure prophylaxis (PrEP), in combination with safer sex practices, to reduce the risk of sexually acquired HIV-1 infection in heterosexual men and women at high risk of acquiring HIV infection who meet both of the following:
 - Condomless vaginal or anal sex; and
 - Ongoing sexual relationship with an HIV-positive partner who is not receiving stable ART and/or does not have an HIV viral load <200 copies/ mL. (i.e. not on ART or >200 copies/mL).

PEOPLE WHO INJECT DRUGS (PWID)

- For pre-exposure prophylaxis (PrEP) for PWID who are at high risk of acquiring HIV infection and meet both of the following:
 - Report sharing of injection equipment; and
 - Have an HIV-positive injecting partner who is not receiving stable ART and/or does not have an HIV viral load < 200 copies/mL.

Clinical Notes:

- PrEP should be part of a combination prevention strategy that includes behavioural interventions such as condoms and risk reduction counseling.
- PrEP is not recommended in the context of a stable closed relationship with a single partner with no or negligible risk of having transmissible HIV.

Note regarding daily versus “on-demand” dosing:

- As stated in the Canadian Guideline, **daily** emtricitabine/tenofovir disoproxil fumarate (TDF/FTC) is currently the PrEP regimen of choice because it has been the most widely evaluated in high quality studies, and “on-demand” dosing is currently an off-label use of TDF/FTC in Canada. The on-demand regimen requires taking the drug 24 hours before sexual activity, every 24 hours during the sexual activity, and 24 hours after the last sexual encounter. A randomized placebo-controlled trial among MSM in France and Montreal found high efficacy among men who had frequent sex and who regularly took an average of 4 pills per week. These results suggest an on-demand strategy may be less effective for MSM who have less frequent sex because consistent pill use is important to achieve high levels of drugs in the body. A subsequent sub-study found that an on-demand strategy (median 9.5 pills/month) remained highly effective for MSM having infrequent sex (median 5x/month). The

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

implication is that on demand' PrEP compared with daily, continuous PrEP may decrease the cost of drugs while preventing similar numbers of infections. However, study of how on-demand PrEP would work in "real life" settings outside of a placebo-controlled trial are required.

***ENCORAFENIB** (*Braftovi 75mg Capsule*)

METASTATIC MELANOMA

- In combination with binimetinib for the treatment of patients with BRAF V600 mutation-positive unresectable or metastatic melanoma.

Clinical Notes:

- Patients should have a good performance status.
- If brain metastases are present, patients should be asymptomatic or have stable symptoms.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.

Claim Notes:

- Encorafenib in combination with binimetinib will not be reimbursed in patients who have progressed on BRAF targeted therapy.
- Requests will be considered for patients who received adjuvant BRAF targeted therapy if disease progression occurred at least 6 months following completion of therapy.

METASTATIC COLORECTAL CANCER

- In combination with panitumumab or cetuximab for the treatment of patients with metastatic colorectal cancer who meet all of the following criteria:
 - Presence of BRAF V600E mutation
 - Disease progression following at least one prior therapy in the metastatic setting
 - No previous treatment with an EGFR inhibitor

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.

Claim Note:

- Encorafenib will not be reimbursed in patients who have progressed on BRAF targeted therapy.

ENTACAPONE (*Comtan 200mg Tablet and generic brands*)

- For the treatment of Parkinson's disease as adjunctive therapy in patients who are not well controlled and are experiencing significant "wearing off" symptoms despite optimal therapy with a levodopa and a decarboxylase inhibitor.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

*ENTRECTINIB (*Rozlytrek 100mg and 200mg Capsule*)

ROS-1 POSITIVE NON-SMALL CELL LUNG CANCER OR METASTATIC NON-SMALL CELL LUNG CANCER

- For the first-line treatment of patients with ROS-1 positive locally advanced (not amenable to curative therapy) or metastatic non-small cell lung cancer (NSCLC).

Clinical Notes:

- Patients should have a good performance status.
- Treatment should continue until disease progression or unacceptable toxicity.

UNRESECTABLE LOCALLY ADVANCED OR METASTATIC EXTRACRANIAL SOLID TUMORS WITH A NTRK GENE FUSION

- For the treatment of adult patients with unresectable locally advanced or metastatic extracranial solid tumors with NTRK gene fusion without a known acquired resistance mutation. Eligible patients are not candidates for surgery and/or radiation due to risk of substantial morbidity and have no satisfactory treatment options.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.
- CNS metastases are stable if present.
- Patients with prior progression on an NTRK inhibitor are not eligible.

*ENZALUTAMIDE (*Xtandi 40mg Capsule*)

METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (mCRPC)

- For the treatment of patients with metastatic castration-resistant prostate cancer (mCRPC).

Clinical Notes:

- Patients should have a good performance status and no risk factors for
- seizures.
- Treatment should continue until disease progression or unacceptable toxicity.
- Requests for enzalutamide will not be considered for patients who experience disease progression on apalutamide or darolutamide.

METASTATIC CASTRATION-SENSITIVE PROSTATE CANCER (mCSPC)

- In combination with androgen deprivation therapy (ADT) for the treatment of patients with metastatic castration-sensitive prostate cancer (mCSPC).

Clinical Notes:

- Patients should have a good performance status and no risk factors for seizures.
- Treatment should continue until disease progression or unacceptable toxicity.
- Patients should have had no prior ADT in the metastatic setting, or are within 6 months of initiating ADT in the metastatic setting with no disease progression.
- Patients will be eligible if they received ADT in the non-metastatic setting as long as at least a one year interval has passed since completion.
- Requests for enzalutamide will not be considered for patients who experience disease progression on apalutamide or darolutamide.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

NON-METASTATIC CASTRATION-RESISTANT PROSTATE CANCER (NMCRPC)

- In combination with androgen deprivation therapy (ADT) for the treatment of patients with non-metastatic castration-resistant prostate cancer (nmCRPC) who have no detectable distant metastases (M0) by either CT, MRI or technetium-99m bone scan and who are at high risk of developing metastases¹.

¹ High risk of developing metastases is defined as a prostate-specific antigen (PSA) doubling time of ≤ 10 months during continuous ADT.

Clinical Notes:

- Patients should have a good performance status and no risk factors for seizures.
- Treatment should continue until radiographic disease progression or unacceptable toxicity.
- Castration-resistance must be demonstrated during continuous ADT and is defined as 3 PSA rises at least one week apart, with the last PSA greater than 2 ng/mL.
- Castrate levels of testosterone must be maintained.
- Patients with N1 disease, pelvic lymph nodes less than 2cm in short axis located below the common iliac vessels are eligible for enzalutamide.
- Enzalutamide will not be funded for patients who experience disease progression on apalutamide or darolutamide.

EPINEPHRINE (*Allerject 0.3mg and 0.15mg Autoinjector, Emerade 0.15mg, 0.3 mg and 0.5mg Prefilled Pen and Epipen 0.15mg and 0.3mg Injection*)

- For the emergency treatment of anaphylactic reactions, when out of reach of immediate medical attention.

Note:

- Regular benefit, but with a quantity limit of two injections per fiscal year. Additional units require an exception status request.

EPLERENONE (*Inspra 25mg, 50mg Tablet and generic brands*)

- For patients >55 years with mild to moderate HF on standard HF treatments with $EF \leq 30\%$ (or $\leq 35\%$ if QRS duration >130ms) and recent (6 months) hospitalization for CV disease or with elevated BNP or NT-proBNP levels.

Notes:

- Requests will be considered from practitioners with a specialty in cardiology.
- Patients must be on optimal therapy with an angiotensin-converting-enzyme (ACE) inhibitor, an angiotensin-receptor blocker (ARB), or both and a beta-blocker (unless contraindicated) at the recommended dose or maximal tolerated dose.

EPTINEZUMAB (*Vyepti 100mg/1mL IV*)

- For the treatment of patients with episodic¹ or chronic migraine², who have experienced an inadequate response, intolerance, or contraindication to at least two oral prophylactic migraine medications of different classes.

Renewal:

- Proof of beneficial clinical effect, defined as a reduction of at least 50% in the average number of migraine days per month at the time of first renewal compared with baseline.
- For subsequent renewals, proof that the initial 50% reduction in the average number of migraine days per month has been maintained.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

- Baseline number of headache and migraine days per month must be provided at the time of initial request.
- ¹Episodic migraine: migraine headaches on at least 4 days per month and less than 15 headache days per month for more than 3 months.
- ²Chronic migraine: headaches for at least 15 days per month for more than 3 months of which at least eight days per month are with migraine.

Claim Notes:

- Initial approval: 6 months
- Renewal approval: 1 year
- Must be prescribed by a physician who has experience in the management of migraine headaches.

***ERYTHROPOIETIN** (*Eprex Multidose Vial and Syringe Injection*)

- For the treatment of transfusion dependent patients with hematologic malignancies who have a baseline anemia of $\leq 90\text{g/L}$ and whose transfusion requirements are ≥ 2 units of packed red blood cells per month over 3 months
- Initial approval for 6 months with the documentation of dose, hemoglobin and therapeutic outcome (number of transfusions).
- Subsequent 6-month approvals are dependent on evidence of satisfactory clinical response or reduced treatment requirement to less than 2 units of PRBC monthly.
- If transfusion requirements increase to ≥ 2 units/ month (over a 3-month period), one dose increase may be attempted (maximum dose 60,000iu per week).

ESLICARBAZEPINE (*Aptiom 200mg, 400mg, 600mg and 800mg Tablet*)

- For the adjunctive treatment of refractory partial-onset seizures (POS) in patients who are currently receiving two or more antiepileptic drugs, and have had an inadequate response or intolerance to at least three other antiepileptic drugs.

Claim Note:

- The patient must be under the care of a physician experienced in the treatment of epilepsy.

***ESTRADIOL** (*Estrogel Topical Gel*)

- For the treatment of menopausal symptoms in women who cannot tolerate the oral forms of hormone replacement therapy.

ETANERCEPT (*Brenzys 50 mg/mL Prefilled Pen and 50 mg/mL Prefilled Syringe; Erelzi 25mg/0.5mL and 50mg/mL Prefilled Syringe and 50mg/mL Prefilled Autoinjector*)**ANKYLOSING SPONDYLITIS**

- For the treatment of patients with moderate to severe ankylosing spondylitis (Bath AS Disease Activity Index (BASDAI) score ≥ 4 on 10 point scale) who:
 - Have axial symptoms¹ and who have failed to respond to the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each, or in whom NSAIDs are contraindicated;
 - OR
 - Have peripheral symptoms and who have failed to respond to, or have contraindications to, the sequential

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each and have had an inadequate response to an optimal dose or maximal tolerated dose of a DMARD.

- Must be prescribed by a rheumatologist or prescriber with a specialty in rheumatology.
- Requests for renewal must include information showing the beneficial effects of the treatment, specifically:
 - A decrease of at least 2 points on the BASDAI scale, compared with the pre-treatment score; OR
 - Patient and expert opinion of an adequate clinical response as indicated by a significant functional improvement (measured by outcomes such as HAQ or "ability to return to work").

Claim Notes:

- Initial coverage period 6 months.
- Maximum dose 50mg per week.
- Concurrent use of biologics not approved.

¹Patients with recurrent uveitis (2 or more episodes within 12 months) as a complication of axial disease do not require a trial of 2 NSAIDs.

POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS

- For the treatment of polyarticular juvenile idiopathic arthritis (pJIA) with the following criteria:
 - For patients aged 4-17 years with moderate or severe pJIA who have had an inadequate response to one or more disease-modifying anti-rheumatic drugs (DMARDs); and
 - Treatment must be initiated by a rheumatologist who is familiar with the use of DMARDs and/or biologic DMARDs in children.

PSORIASIS

- For patients with severe, debilitating chronic plaque psoriasis who meet all of the following:
 - Body surface area (BSA) involvement of >10% and/or significant involvement of the face, hands, feet or genitals;
 - Failure to, contraindication to or intolerant of methotrexate and cyclosporine;
 - Failure to, intolerant of or unable to access phototherapy;
 - Written request of a dermatologist or prescriber with a specialty in dermatology.
- Continued coverage is dependent on evidence of improvement, specifically:
 - A ≥75% reduction in the Psoriasis Area and Severity Index (PASI) score; or
 - A ≥50% reduction in PASI with a ≥5-point improvement in DLQI (Dermatology Life Quality Index); or
 - Significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals.

Clinical Note:

- Treatment should be discontinued if a response has not been demonstrated after 12 weeks.

Claim Notes:

- Concurrent use of biologics not approved.
- Initial approval 12 weeks.
- Maximum dose 50mg biweekly for initial, then 50mg per week.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

PSORIATIC ARTHRITIS

- For the treatment of patients with predominantly axial psoriatic arthritis who are refractory, intolerant or have contraindications to the sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each.
- For the treatment of patients with predominantly peripheral psoriatic arthritis who are refractory, intolerant or have contraindications to:
 - The sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each;
AND
 - Methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age) for a minimum of 8 weeks;
AND
 - Leflunomide for a minimum of 10 weeks or sulfasalazine for a minimum of 3 months.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- 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 to treatments. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial approval 3 months.
- Maximum dose of 50mg per week.
- Renewal approval: 1 year. Confirmation of continued response required.

RHEUMATOID ARTHRITIS

- For the treatment of severely active rheumatoid arthritis, in combination with methotrexate or other DMARDs, in adult patients who are refractory or intolerant to:
 - Methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age), or use in combination with another DMARD, for a minimum of 12 weeks
AND
 - Methotrexate in combination with at least two other DMARDs, such as hydroxychloroquine and sulfasalazine, for a minimum of 12 weeks.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- Optimal treatment response to DMARDs may take up to 24 weeks, however coverage of a biologic therapy can be considered if no improvement is seen after 12 weeks of triple DMARD use.
- If patient factors (e.g. intolerance) prevent the use of triple DMARD therapy, these must be described and dual therapy with DMARDs must be tried.
- Refractory is defined as lack of effect at the recommended doses and for duration of treatments specified above.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Intolerant is defined as demonstrating serious adverse effects or contraindications to treatments as defined in product monographs. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial Approval: 6 months.
- Renewal Approval: 1 year. Confirmation of continued response is required.
- Maximum Dosage Approved: 25mg twice a week or 50mg once a week with no dose escalation permitted.

***EVEROLIMUS (Afinitor 2.5mg, 5mg, and 10mg Tablet and generic brands)**

METASTATIC RENAL CELL CARCINOMA (RCC)

- For the treatment of patients with advanced or metastatic renal cell carcinoma following disease progression on tyrosine kinase inhibitor therapy.

Clinical Notes:

- Patients must have a good performance status.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.
- Requests for everolimus will not be considered for patients who experience disease progression on axitinib, cabozantinib or nivolumab monotherapy.

HORMONE RECEPTOR POSITIVE, HER2 NEGATIVE-ADVANCED BREAST CANCER

- In combination with exemestane for postmenopausal patients (ECOG PS ≤ 2) with documented hormone receptor positive, HER2 negative-advanced breast cancer after recurrence or progression following a non-steroidal aromatase inhibitor (NSAI).

Note:

- It may be clinically reasonable to use the combination in patients with treated and stable brain metastasis.

METASTATIC PANCREATIC NEUROENDOCRINE TUMORS (PNET)

- For the treatment of patients with progressive, unresectable, well or moderately differentiated, locally advanced or metastatic pancreatic neuroendocrine tumors (pNET) with good performance status (ECOG 0-2), until disease progression.

Note:

- Patients whose disease progresses on sunitinib are not eligible for funded treatment with everolimus for pNET.

NEUROENDOCRINE TUMOURS OF GASTROINTESTINAL OR LUNG ORIGIN

- As a single agent treatment for patients with unresectable, locally advanced or metastatic; well-differentiated non-functional neuroendocrine tumours (NETs) of gastrointestinal or lung origin (GIL) in adults with documented radiological disease progression within six months and with a good performance status. Treatment should continue until confirmed disease progression or unacceptable toxicity.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

EVOLOCUMAB (*Repatha 140mg/mL Prefilled Syringe and 120mg/mL Automated Mini Doser*)

- For the treatment of heterozygous familial hypercholesterolemia (HeFH) in adult patients who require additional lowering of low-density lipoprotein cholesterol (LDL-C) if the following criteria are met:
 - Definite or probable diagnosis of HeFH using the Simon Broome or Dutch Lipid Network criteria or genetic testing; and
 - Patient is unable to reach LDL-C target (less than 2.0 mmol/L or at least a 50% reduction in LDL-C from untreated baseline) despite confirmed adherence to at least 3 months of continuous treatment with:
 - high-dose statin (e.g., atorvastatin 80 mg, rosuvastatin 40 mg) in combination with ezetimibe; or
 - ezetimibe alone if high dose statin is not possible due to rhabdomyolysis, contraindication or intolerance
- **Initial renewal criteria:**
 - A reduction in LDL-C of at least 40% from baseline or has reached a target LDL-C less than 2.0 mmol/L.
- **Subsequent renewal criteria:**
 - The patient continues to maintain a reduction in LDL- C of at least 40% from baseline or has reached a target LDL-C less than 2.0 mmol/L.

Clinical Notes:

- LDL-C levels must be provided.
- Intolerance to high dose statin will be considered if patient has developed documented, myopathy or abnormal biomarkers (i.e. creatinine kinase greater than 5 times the upper limit of normal) after trial of at least two statins and
 - for each statin, dose reduction was attempted rather than statin discontinuation, and intolerance was reversible upon statin discontinuation, but reoccurred with statin re-challenge where clinically appropriate; and
 - at least one statin was initiated at the lowest daily starting dose; and
 - other known causes of intolerance or abnormal biomarkers have been ruled out.
- For patients who cannot take a statin due to an intolerance or contraindication, details must be provided (ie. confirmed rhabdomyolysis, active liver disease, unexplained persistent elevations of serum transaminases exceeding three times the upper limit of normal).
- For patients who cannot take ezetimibe due to an intolerance or contraindication, details must be provided.

Claim Notes:

- Maximum dose approved: 140mg every 2 weeks or 420mg monthly
- Initial approval: 6 months
- Renewal approval: 1 year

FEBUXOSTAT (*Uloric 80mg Tablet and generic brands*)

- For the treatment of symptomatic gout in patients who have documented hypersensitivity to allopurinol.

***FEDRATINIB** (*Inrebic 100mg Capsule*)

- For the treatment of splenomegaly and/or disease-related symptoms in adult patients with intermediate-2 or high-risk primary myelofibrosis, post-polycythemia vera myelofibrosis, or post-essential thrombocythemia myelofibrosis, who have a contraindication or intolerance to ruxolitinib.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.

Claim Note:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00904799

***FENTANYL** (*Fentanyl 12mcg/hr, 25mcg/hr, 50mcg/hr, 75mcg/hr, 100mcg/hr Transdermal System and generic brands*)

- For the treatment of malignant or chronic non-malignant pain in adult patients who were previously receiving continuous opioid administration (i.e., not opioid naive), or who are unable to take oral therapy.

FESOTERODINE FUMARATE (*Toviaz 4mg and 8mg Tablet and generic brands*)

- See [OAB Medications](#)

***FERRIC DERISOMALTOSE** (*Monoferric 100 mg/mL IV Injection*)

- For the treatment of iron deficiency anemia in patients who:
 - are intolerant to oral iron replacement products,
OR
 - have not responded to an adequate trial of oral iron

Notes:

- Given the safety concerns associated with IV iron, it is expected that the patients will be carefully screened and will have tried various oral iron options before being eligible for IV iron.
- Details regarding oral iron tried, length of therapy, and outcome must be provided.

FIDAXOMICIN (*Dificid 200mg Tablet*)

- For the treatment of Clostridium Difficile Infection (CDI) where the patient:
 - has experienced a third or subsequent episode within 6 months of treatment with vancomycin for prior episode(s), with no previous trial of fidaxomicin; OR
 - has experienced treatment failure¹ with oral vancomycin for the current CDI episode; OR
 - has had a documented allergy (immune-mediated reaction) to oral vancomycin; OR
 - has experienced a severe adverse reaction or intolerance² to oral vancomycin treatment that resulted in the discontinuation of vancomycin therapy.
- **Re-treatment criteria:**
 - Re-treatment with fidaxomicin will only be considered for an early relapse occurring within 30 days of the completion of the most recent fidaxomicin course.
 - Relapse/recurrence occurring beyond 30 days after the completion of the most recent fidaxomicin course will require a trial with vancomycin, unless there is a documented allergy, severe adverse reaction or intolerance to prior oral vancomycin use.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

1. Treatment failure is defined as 7 days of vancomycin therapy without acceptable clinical improvement.
2. Details of severe adverse reaction or intolerance must be provided and should be clinically related to oral administration of vancomycin.

Claim Note:

- Requests will be approved for 200mg twice a day for 10 days.

FINERENONE (*Kerendia 10mg and 20mg Tab*)

- For the treatment of patients with chronic kidney disease (CKD) and type 2 diabetes (T2D) who have an estimated glomerular filtration rate (eGFR) level of at least 25 mL/min/1.73 m² and albuminuria level of at least 30 mg/g (or 3 mg/mmol).

Exclusion Criteria:

- Patients with chronic heart failure (CHF) New York Heart Association (NYHA) class II to IV; OR
- Patients receiving a mineralocorticoid receptor antagonist (MRA).

Discontinuation Criteria:

- eGFR less than 15 mL/min/1.73 m²; OR
- Urinary albumin-to-creatinine ratio (UACR) increased from baseline level.

Claim Notes:

- Must be prescribed by, or in consultation with, a nephrologist with experience in the diagnosis and management of patients with CKD and T2D.
- Approval: 1 year

FINGOLIMOD (*Gilenya 0.5mg Capsule and generic brands*)

- For the treatment of patients with relapsing remitting multiple sclerosis (RRMS) who meet **all** of the following criteria:
 - have failed to respond to a full and adequate course¹ of at least one disease modifying therapy (DMT) publicly insured in Nova Scotia as an initial therapy, or has contraindications/intolerance² to at least two initial therapies;
 - one or more clinically disabling relapses in the previous year;
 - significant increase in T2 lesion load compared with that from a previous MRI scan (i.e. 3 or more new lesions) or at least one gadolinium-enhancing lesion;
 - requested and followed by a neurologist experienced in the management of RRMS;
 - recent expanded disability status scale (EDSS) score of 5.5 or less (i.e. patients must be able to ambulate at least 100 meters without assistance).
- Dosage: 0.5mg daily
- Approval period: 1 year
- **Exclusions:**
 - not funded in combination with other disease modifying therapies;
 - not funded in patients with an EDSS>5.5;
 - not funded in patients who have had a heart attack or stroke in the last six months of funding request, patients with a history of sick sinus syndrome, atrioventricular block, significant QT prolongations, bradycardia, ischemic heart disease, or congestive heart failure;

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- not funded in patients <18 years of age;
- not funded due to needle phobia or preference for oral therapy over injection in patients without clinical contraindications to interferon or glatiramer therapy.

Note:

- Skin reactions at the site of injection do not qualify as contraindications to interferon or glatiramer therapy.

Renewal:

- EDSS score ≤ 5.5 (i.e. patients must be able to ambulate at least 100 meters without assistance). Date and details of the most recent neurological examination and EDSS scores must be provided (exam must have occurred within that last 90 days); AND
- Patients must be stable or have experienced no more than 1 disabling attack/relapse in the past year.

Of Note:

1. Failure to respond to full and adequate courses: defined as a trial of at least 6 months of interferon or glatiramer therapy AND experienced at least one disabling relapse (attack) while on interferon or glatiramer therapy
2. Intolerance is defined as: documented serious adverse effects or contraindications that are incompatible with further use of that class of drug

***FLUCONAZOLE** (*Diflucan POS 10mg/mL*)

- For the treatment of oropharyngeal candidiasis when nystatin has failed, or for systemic infections when oral tablets are not an option.

***FLUDARABINE** (*Fludara 10mg Tablet*)

- For the treatment of chronic lymphocytic leukemia (CLL), in patients with an ECOG performance status of 0-2, when:
 - the patient has failed to respond or relapsed during or after previous therapy with an alkylating agent, and
 - intravenous administration is not desirable.

FLUOROQUINOLONES, OPHTHALMIC (*Ciprofloxacin, Gatifloxacin, Moxifloxacin and Ofloxacin*)

- For the treatment of eye infections upon the order of an ophthalmologist, ophthalmology resident, prescribing optometrist or other prescriber who has a specialty in ophthalmology. **[Criteria Code 01]**

***FLUOROQUINOLONES, ORAL** (*Ciprofloxacin, Norfloxacin*)

- For the treatment of patients intolerant or allergic (hypersensitivity reaction) to all other effective oral agents. **[Criteria Code 01]**
- For the treatment of aerobic, gram-negative infections which are resistant to other suitable oral agents. **[Criteria Code 02]**
- For the oral treatment of multi-resistant, aerobic, gram-negative infections traditionally requiring parenteral therapy (e.g., osteomyelitis, complicated urinary tract infections, bacterial pneumonia in cystic fibrosis, prostatitis) for which other oral agents are not effective or available. **[Criteria Code 03]**
- For infections due to *Pseudomonas aeruginosa* (ciprofloxacin is the preferred agent). **[Criteria Code 04]**
- For the treatment of necrotizing (malignant) otitis externa. **[Criteria Code 05]**
- For the prevention of endophthalmitis in patients who have had cataract surgery involving an unplanned vitrectomy (ciprofloxacin). **[Criteria Code 06]**
- For prevention of recurrent spontaneous bacterial peritonitis (norfloxacin). **[Criteria Code 07]**

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

*FLUOROQUINOLONES, RESPIRATORY (*Levofloxacin, Moxifloxacin*)

- for the completion of therapy instituted in the hospital setting for the treatment of nosocomial pneumonia, community acquired pneumonia (CAP) or acute exacerbation of chronic bronchitis (AECB). **[Criteria Code 01]**
 - for the treatment of severe pneumonia in nursing home patients. **[Criteria Code 02]**
 - for the treatment¹ of CAP in patients with comorbidity² upon radiographic confirmation of pneumonia, or who have failed first line therapies (macrolide, doxycycline, amoxicillin-clavulanate). **[Criteria Code 03]**
 - for the treatment¹ of AECB in complicated patients³ who have failed treatment with one of the following: amoxicillin, doxycycline, TMP-SMX, cefuroxime, macrolide, ketolide or amoxicillin-clavulanate. **[Criteria Code 04]**
1. If treated with an antibiotic within the past 3 months choose an antibiotic from a different class.
 2. Comorbidity includes chronic lung disease, malignancy, diabetes, liver failure, renal failure, congestive heart failure, use of antibiotics or steroids in the past 3 months, suspected macroaspiration, hospitalization within last 3 months, HIV/AIDS, smoking, malnutrition or acute weight loss.
 3. Complicated AECB defined as increased cough and sputum, sputum purulence and increased dyspnea and FEV1 < 50% predicted or FEV1 50% - 65% and one of the following: ≥4 exacerbations per year, ischemic heart disease, chronic oral steroid use or antibiotic use in past 3 months

*FLUOXETINE (*Prozac 20mg/5mL Syr and generic brands*)

- For use in patients for whom oral capsules are not an option.

FLUTICASONE FUROATE AND VILANTEROL (AS TRIFENATATE) (*Breo Ellipta 100mcg/25mcg and 200mcg/25mcg Powder for Inhalation*)

- See [Long-Acting Beta₂-Agonists/Inhaled Corticosteroids](#)

FLUTICASONE FUROATE/UMECLIDINIUM/VILANTEROL (*Trelegy Ellipta 100mcg/62.5mcg/25mcg*)

- For the treatment of chronic obstructive pulmonary disease (COPD), as defined by spirometry, in patients who experience inadequate control while being treated with a long-acting beta-2 agonist/long-acting muscarinic antagonist (LABA/LAMA).

Clinical Notes:

- COPD is defined by spirometry as a post-bronchodilator FEV₁/FVC ratio of less than 0.70. Spirometry reports from any point in time will be accepted. If spirometry cannot be obtained, reasons must be clearly explained and other evidence of COPD severity provided (i.e. MRC Dyspnea Scale Score grade).
- Inadequate control while being treated with a LABA/LAMA for at least two months is defined as persistent symptoms or experiencing two or more exacerbations of COPD in the previous year requiring treatment with antibiotics and/or systemic corticosteroids or at least one exacerbation of COPD requiring hospitalization.
- Patients should not be started on a LABA, LAMA and an inhaled corticosteroid (triple inhaled therapy) as initial therapy.

FORMOTEROL (*Oxeze 6mcg/Dose, 12mcg/Dose Turbuhaler*)

- See [Long-Acting Beta₂-Agonists](#)

FORMOTEROL, IN COMBINATION (*Symbicort 100/6mcg, 200/6mcg Turbuhaler and Zenhale 50/5mcg, 100/5mcg, 200/5mcg Inh*)

- See [Long-Acting Beta₂-Agonists/Inhaled Corticosteroids](#)

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FOSFOMYCIN TROMETHAMINE (*Monurol 3g/sachet and generic brands*)

- For the treatment of uncomplicated urinary tract infections in adult female patients where:
 - the infecting organism is resistant to other oral agents [**Criteria Code 01**]; OR
 - other less costly treatments are not tolerated. [**Criteria Code 02**]

FREMANEZUMAB (*Ajovy 225 mg/1.5 mL Prefilled Syringe and 225 mg/1.5 mL Autoinjector*)

- For the treatment of patients with episodic¹ or chronic migraine², who have experienced an inadequate response, intolerance, or contraindication to at least two oral prophylactic migraine medications of different classes.

Renewal:

- Proof of beneficial clinical effect, defined as a reduction of at least 50% in the average number of migraine days per month at the time of first renewal compared with baseline
- For subsequent renewals, proof that the initial 50% reduction in the average number of migraine days per month has been maintained

Clinical Notes:

- Baseline number of headache and migraine days per month must be provided at the time of initial request.
- ¹ Episodic migraine: migraine headaches on at least 4 days per month and less than 15 headache days per month for more than 3 months
- ² Chronic migraine: headaches for at least 15 days per month for more than 3 months of which at least eight days per month are with migraine.

Claim Notes:

- Approvals: 6 months
- Renewal approval: 1 year
- Must be prescribed by a physician who has experience in the management of migraine headaches.

GALCANEZUMAB (*Emgality 120mg/mL Prefilled Pen and 120mg/mL Prefilled Syringe*)

Initiation:

- For the treatment of patients with episodic¹ or chronic migraine², who have experienced an inadequate response, intolerance, or contraindication to at least two oral prophylactic migraine medications of different classes.

Renewal:

- Proof of beneficial clinical effect, defined as a reduction of at least 50% in the average number of migraine days per month at the time of first renewal compared with baseline.
- For subsequent renewals, proof that the initial 50% reduction in the average number of migraine days per month has been maintained.

Clinical Notes:

- Baseline number of headache and migraine days per month must be provided at the time of initial request.
- ¹Episodic migraine: migraine headaches on at least 4 days per month and less than 15 headache days per month for more than 3 months.
- ²Chronic migraine: headaches for at least 15 days per month for more than 3 months of which at least eight days per month are with migraine.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Notes:

- Initial approval: 6 months
- Renewal approval: 1 year
- Must be prescribed by a physician who has experience in the management of migraine headaches.

GATIFLOXACIN (*Zymar 0.3% Ophthalmic Solution and generic brands*)

- See [Fluoroquinolones, Ophthalmic](#)

***GILTERITINIB** (*Xospata 40mg Tablet*)

- As monotherapy for the treatment of adult patients with relapsed or refractory FMS-like tyrosine kinase 3 (FLT3)-mutated acute myeloid leukemia (AML) who meet the following criteria:
 - Confirmed positive for FLT3 mutation at the time of relapse or determination of refractory disease, eligible FLT3 mutations include FLT3-ITD, and FLT3-TKD.

Clinical Notes:

- Patients should have a good performance status.
- Treatment with gilteritinib should be continued as long as clinical benefit is observed, or until disease progression or unacceptable toxicity, whichever occurs first.
- Patients previously treated with midostaurin are eligible for gilteritinib provided all other criteria are met.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904658
 - 00904659

GIVOSIRAN (*Givlaari 189mg/mL Vial*)

- For the treatment of acute hepatic porphyria (AHP) in adults:

Initiation:

- Reimbursement of givosiran should be restricted to patients with 4 or more attacks requiring either hospitalization, an urgent health care visit, or IV hemin in the year before the prescribing date.

Renewal:

- A reduction in the annualized attack rate after 12 months of therapy compared to baseline.

Claim Notes:

- Prescription should be restricted to a clinician experienced in the management of AHP.
- Should not be used in combination with prophylactic hemin.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and the following PINs:
 - 00904990
 - 00904991
 - 00904992
 - 00904993
 - 00904994
 - 00904995

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

GLATIRAMER ACETATE (*Copaxone 20mg/mL Syringe Injection*)

For glatiramer acetate-naïve patients whose glatiramer acetate therapy is initiated after April 1, 2020, the Glatect brand will be the product approved.

Prescribed by a neurologist with experience in the treatment of multiple sclerosis for patients who meet the following criteria:

Treatment initiation:

- Diagnosis of Multiple Sclerosis with a relapsing course*:
 - Includes relapsing-remitting MS and secondary progressive MS with clear superimposed relapses;
 - Does not include primary progressive MS, progressive- relapsing or secondary progressive MS without relapses;
and
 - Disability judged to be equivalent to Expanded Disability Status Score (EDSS) of 5.5 or less (exceptions are permitted in special cases).

Renewal:

- EDSS not greater than 6.0 for at least 12 months in the absence of relapses.
- Patients must be assessed for compliance and for any therapy related side effects that are intolerable.

Exclusions:

- Concurrent illness likely to alter compliance or substantially reduce life expectancy

* Relapsing course is defined as evidence of one relapse in the past 18 months or two relapses in the past 3 years.

GLATIRAMER ACETATE (*Glatect 20mg Pre-Filled Syringe*)

For glatiramer acetate-naïve patients whose glatiramer acetate therapy is initiated after April 1, 2020, the Glatect brand will be the product approved.

Prescribed by a neurologist with experience in the treatment of multiple sclerosis for patients who meet the following criteria:

Treatment initiation:

- Diagnosis of Multiple Sclerosis with a relapsing course:
 - Includes relapsing-remitting MS and secondary progressive MS with clear superimposed relapses;
 - Does not include primary progressive MS, progressive- relapsing or secondary progressive MS without relapses;
and
 - Disability judged to be equivalent to Expanded Disability Status Score (EDSS) of 5.5 or less (exceptions are permitted in special cases).

Renewal:

- EDSS not greater than 6.0 for at least 12 months in the absence of relapses.
- Patients must be assessed for compliance and for any therapy related side effects that are intolerable.

Exclusions:

- Concurrent illness likely to alter compliance or substantially reduce life expectancy

* Relapsing course is defined as evidence of one relapse in the past 18 months or two relapses in the past 3 years.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

GLECAPREVIR/ PIBRENTASVIR (*Maviret 100mg/40mg Tablet*)

- For treatment-naïve or treatment-experienced adult patients with chronic hepatitis C virus (HCV) who meet the following criteria:

	Approval Period and Regimen
Genotypes 1, 2, 3, 4, 5 or 6 <ul style="list-style-type: none">Treatment-naïve	<ul style="list-style-type: none">8 weeks
Genotypes 1, 2, 4, 5 or 6 <ul style="list-style-type: none">Treatment-experienced with regimens containing peginterferon/ribavirin (PR) and/or sofosbuvir (SOF)	<ul style="list-style-type: none">8 weeks (12 weeks with cirrhosis)
Genotype 1 <ul style="list-style-type: none">NS5A inhibitor treatment-naïve and treatment-experienced with regimens containing:<ul style="list-style-type: none">Boceprevir/PR; orSimeprevir (SMV)/SOF; orSMV/PR; orTelaprevir/PR	<ul style="list-style-type: none">12 weeks
Genotype 1 <ul style="list-style-type: none">NS3/4A inhibitor treatment-naïve and treatment-experienced with regimens containing:<ul style="list-style-type: none">Daclatasvir (DCV)/SOF; orDCV/PR; orLedipasvir/SOF	<ul style="list-style-type: none">16 weeks
Genotype 3 <ul style="list-style-type: none">Treatment-experienced with regimens containing PR and/or SOF	<ul style="list-style-type: none">16 weeks

- The following information is also required:
 - Lab-confirmed hepatitis C genotype 1, 2, 3, 4, 5 or 6
 - Quantitative HCV RNA value within the last 6 months
 - Fibrosis stage

Clinical Note:

- Acceptable methods for the measurement of fibrosis score include Fibrotest, liver biopsy, transient elastography (FibroScan®), serum biomarker panels (such as AST-to-Platelet Ratio Index or Fibrosis-4 score) either alone or in combination.

Claim Notes:

- Must be prescribed by a hepatologist, gastroenterologist, or infectious disease specialist (or other physician experienced in treating a patient with hepatitis C infection).
- Claims will be limited to a 28-day supply.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate

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transactions using the DIN first and then the following PINs:

- 00904394
- 00904395

** [Criteria Code 34] has been added for use effective December 1, 2019 to allow payment of a patient's initial 28 day supply only. Criteria code 34 should be provided by the prescribing physician only.

GLUCAGON (*Baqsimi 3mg Nasal Powder*)

- For the emergency treatment of severe hypoglycemia (SH) reactions for patients who are receiving insulin and at high risk for SH, when impaired consciousness precludes oral carbohydrate.

Claim Note:

- Regular benefit, but with a quantity limit of two (2) devices per fiscal year. Prescribers can submit a request for consideration or contact the Pharmacare Office should beneficiaries require more than two (2) devices per fiscal year (e.g., need for frequent use).

GLYCEROL PHENYLBUTYRATE (*Ravicti 1.1g/mL Oral Liquid*)

- For the chronic management of patients with urea cycle disorders (UCDs).

Clinical Note:

- Diagnosis must be confirmed by blood, enzymatic, biochemical or genetic testing.

Claim Notes:

- Must be prescribed by, or in consultation with, a prescriber experienced in the treatment of UCDs.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904360
 - 00904361

GLYCOPYRRONIUM BROMIDE (*Seebri 50mcg Capsule for Inhalation*)

- See [Long-Acting Muscarinic Antagonists](#)

GOLIMUMAB (*Simponi 50mg/0.5ml and 100mg/1mL Autoinjector and Prefilled Syringe*)

ANKYLOSING SPONDYLITIS

- For the treatment of patients with moderate to severe ankylosing spondylitis (Bath AS Disease Activity Index (BASDAI) score ≥ 4 on 10 point scale) who:
 - Have axial symptoms¹ and who have failed to respond to the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each, or in whom NSAIDs are contraindicated;
 - OR
 - Have peripheral symptoms and who have failed to respond to, or have contraindications to, the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each and have had an inadequate response to an optimal dose or maximal tolerated dose of a DMARD.
- Must be prescribed by a rheumatologist or prescriber with a specialty in rheumatology.
- Requests for renewal must include information showing the beneficial effects of the treatment, specifically:
 - A decrease of at least 2 points on the BASDAI scale, compared with the pre-treatment score; OR

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- Patient and expert opinion of an adequate clinical response as indicated by a significant functional improvement (measured by outcomes such as HAQ or "ability to return to work").

Claim Notes:

- Initial period 16 weeks.
- Maximum dose 50mg per month.
- Concurrent use of biologics not approved.

1. Patients with recurrent uveitis (2 or more episodes within 12 months) as a complication of axial disease, do not require a trial of 2 NSAIDs.

PSORIATIC ARTHRITIS

- For the treatment of patients with predominantly axial psoriatic arthritis who are refractory, intolerant or have contraindications to the sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each.
- For the treatment of patients with predominantly peripheral psoriatic arthritis who are refractory, intolerant or have contraindications to:
 - The sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each; AND
 - Methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age) for a minimum of 8 weeks; AND
 - Leflunomide for a minimum of 10 weeks or sulfasalazine for a minimum of 3 months.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- 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 to treatments. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial period 3 months
- Maximum dose 50mg per month.
- Renewal approval: 1 year. Confirmation of continued response required.

RHEUMATOID ARTHRITIS

- For the treatment of severely active rheumatoid arthritis, in combination with methotrexate or other disease-modifying antirheumatic drugs (DMARDs), in adult patients who are refractory or intolerant to:
 - methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age), or use in combination with another DMARD, for a minimum of 12 weeks; AND
 - methotrexate in combination with at least two other DMARDs, such as hydroxychloroquine and

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sulfasalazine, for a minimum of 12 weeks.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- Optimal treatment response to DMARDs may take up to 24 weeks, however coverage of a biologic therapy can be considered if no improvement is seen after 12 weeks of triple DMARD use.
- If patient factors (e.g. intolerance) prevent the use of triple DMARD therapy, these must be described and dual therapy with DMARDs must be tried.
- 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. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial Approval: 6 months.
- Maximum dose 50mg once a month with no escalation permitted.
- Renewal Approval: 1 year. Confirmation of continued response is required.

ULCERATIVE COLITIS

- For the treatment of patients with moderately to severely active ulcerative colitis who have a partial Mayo score > 4, and a rectal bleeding subscore ≥ 2 and are:
 - refractory or intolerant to conventional therapy (i.e. 5-ASA for a minimum of 4 weeks, and prednisone ≥ 40 mg daily for two weeks or IV equivalent for one week); OR
 - corticosteroid dependent (i.e. cannot be tapered from corticosteroids without disease recurrence; or have relapsed within three months of stopping corticosteroids; or require two or more courses of corticosteroids within one year.)
- Renewal requests must include information demonstrating the beneficial effects of the treatment, specifically:
 - a decrease in the partial Mayo score ≥ 2 from baseline, AND
 - a decrease in the rectal bleeding subscore ≥ 1 .

Clinical Notes:

- 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. The nature of intolerance(s) must be clearly documented.
- Patients with severe disease do not require a trial of 5-ASA.

Claim Notes:

- Must be prescribed by a gastroenterologist or physician with a specialty in gastroenterology.
- Concurrent use of biologics not approved.
- Initial Approval: 16 weeks.
- Maximum dose 50mg once a month with no escalation permitted.
- Renewal Approval: 1 year.

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*GRANISETRON (*Kytril 1mg Tablet and generic brands*)

- See [Serotonin \(5-HT₃\) Antagonists](#)

Note:

- Recommended dose is 2mg orally 1 hour pre-chemotherapy or 1mg 1 hour pre-chemotherapy and 1mg 12 hours post-chemotherapy.

GRASS POLLEN ALLERGEN EXTRACT (*Oralair 100 and 300 Unit IR S/L Tablet*)

- For the seasonal treatment of grass pollen allergic rhinitis in patients that have not adequately responded to, or tolerated, conventional pharmacotherapy.

Notes:

- Treatment with 5-GPAE must be prescribed and initiated by physicians with adequate training and experience in the treatment of respiratory allergic diseases.
- Treatment should be initiated four (4) months before onset of pollen season and should only be continued until the end of the season.
- Treatment should not be taken for more than three (3) consecutive years.

GUSELKUMAB (*Tremfya 100mg/mL Autoinjector and Prefilled Syringe*)

PSORIASIS

- For patients with severe, debilitating chronic plaque psoriasis who meet all of the following:
 - Body surface area (BSA) involvement of >10% and/or significant involvement of the face, hands, feet or genitals;
 - Failure to, contraindication to or intolerant of methotrexate and cyclosporine;
 - Failure to, intolerant of or unable to access phototherapy;
 - Written request of a dermatologist or prescriber with a specialty in dermatology.
- Continued coverage is dependent on evidence of improvement, specifically:
 - A >75% reduction in the Psoriasis Area and Severity Index (PASI) score; or
 - A >50% reduction in PASI with a > 5 point improvement in DLQI (Dermatology Life Quality Index); or
 - Significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals.

Clinical Note:

- Treatment should be discontinued if a response has not been demonstrated after 16 weeks.

Claim Notes:

- Concurrent use of biologics not approved.
- Approvals will be for 100mg by subcutaneous injection at weeks 0, 4, followed by maintenance dosing of 100mg every 8 weeks.
- Initial approval period: 16 weeks
- Renewal approval period: 1 year

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

PSORIATIC ARTHRITIS

- For the treatment of patients with predominantly axial psoriatic arthritis who are refractory, intolerant or have contraindications to the sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each.
- For the treatment of patients with predominantly peripheral psoriatic arthritis who are refractory, intolerant or have contraindications to:
 - The sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each; AND
 - Methotrexate (oral or parenteral) at a dose of $\geq 20\text{mg}$ weekly ($\geq 15\text{mg}$ if patient is ≥ 65 years of age) for a minimum of 8 weeks; AND
 - Leflunomide for a minimum of 10 weeks or sulfasalazine for a minimum of 3 months.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- 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 to treatments. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Combined use of more than one biologic DMARD will not be reimbursed.
- Initial approval: 12 weeks, loading dose of 100mg at weeks 0, 4, and 8 weeks
- Maximum dose of 100mg every 8 weeks
- Renewal approval: 1 year. Confirmation of continued response required.

HALOBETASOL PROPIONATE AND TAZAROTENE (*Duobrii 0.01%/0.045% Topical Lotion*)

- Patients must have a clinical diagnosis of moderate to severe plaque psoriasis and an inadequate response to a topical high-potency corticosteroid.

HYDROXYZINE (*10mg, 25mg, 50mg Capsule, generic brands and Atarax Syrup*)

- For chronic urticaria, defined as the presence of hives or lesions for longer than six weeks, which has responded to treatment with hydroxyzine

HUMAN INSULIN R (*Entuzity 500 U/mL KwikPen*)

- For the treatment of patients with diabetes mellitus with unacceptable glycemic control who require more than 200 units of insulin per day, with or without other therapies.

Claims Notes:

- Treatment must be initiated by an endocrinologist or prescriber with a specialty in endocrinology.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

*IBRUTINIB (*Imbruvica 140mg Capsule*)

FIRST LINE CHRONIC LYMPHOCYTIC LEUKEMIA/ SMALL LYMPHOCYTIC LYMPHOMA

- As a single agent treatment option for patients with previously untreated chronic lymphocytic leukemia (CLL)/ small lymphocytic lymphoma (SLL) for whom fludarabine –based treatment is considered inappropriate due to high risk of relapse or refractory disease based on prognostic biomarkers. Treatment should be discontinued upon disease progression or unacceptable toxicity.

Clinical Notes:

- High risk for relapse or refractory disease includes 17p deletion, TP53 mutation, 11q deletion and unmutated IGHV.
- Sequential use of ibrutinib and idelalisib will not be funded, except as a bridge to transplant. Exceptions may be considered in the case of intolerance without disease progression.

RELAPSED/REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA OR SMALL LYMPHOCYTIC LYMPHOMA

- As a treatment option for patients with relapsed and/or refractory chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) who have received at least one prior therapy and are considered inappropriate for treatment or retreatment with a fludarabine-based regimen, including:
 - Patients who received prior fludarabine-based treatment and had a progression free interval of less than three years
 - Patients who received prior fludarabine-based treatment and had a progression free interval of greater than three years, but are now considered unfit for fludarabine-based retreatment due to age ≥ 70 , or age ≥ 65 and the presence of comorbidities (Cumulative Illness Rating Scale [CIRS] ≥ 6 or creatinine clearance <70 ml/min)
 - Patients who did not receive prior fludarabine-based treatment because they were considered unfit, and who relapsed after at least two cycles of alkylator-based therapy, regardless of the progression free interval after that therapy

RELAPSED/REFRACTORY MANTLE CELL LYMPHOMA

- As a single agent treatment option for patients with relapsed or refractory mantle cell lymphoma who have received at least one prior therapy. Patients should have a good performance status. Treatment should be discontinued upon disease progression or unacceptable toxicity.

ICATIBANT (*Firazyr 30mg/30mL Prefilled Syringe*)

- For the treatment of acute attacks of hereditary angioedema (HAE) in adults with lab confirmed c1-esterase inhibitor deficiency (type I or type II) under the following conditions:
 - treatment of non-laryngeal attacks of at least moderate severity, or
 - treatment of acute laryngeal attacks

Notes:

- Limited to a single dose for self-administration per attack
- Be prescribed by physicians with experience in the treatment of HAE

Claim Note:

- Maximum of two doses on hand at any time.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

***IDELALISIB** (*Zydelig 100mg and 150mg Tablet*)

- In combination with rituximab for the treatment of patients with relapsed chronic lymphocytic leukemia (CLL). Treatment should continue until unacceptable toxicity or disease progression

***IMIQUIMOD** (*Aldara P 5% Cream and generic brands*)

- For the treatment of external genital and perianal warts and condyloma acuminata in adults.
- For the treatment of actinic keratosis on the head and neck in patients who have failed treatment with 5FU and cryotherapy.
- For the treatment of biopsy-confirmed primary superficial basal cell carcinoma:
 - with a tumor diameter of ≤ 2 cm; AND
 - located on the trunk, neck or extremities (excluding hands and feet); AND
 - where surgery or eradication therapy is not medically indicated;
 - Recurrent lesions in previously irradiated area; OR
 - Multiple lesions, too numerous to irradiate or remove surgically.
 - approval period: 6 weeks

Note:

- Surgical management should be considered first-line for superficial basal cell carcinoma in most patients, especially for isolated lesions.

INCObOTULINUMTOXIN-A (*Xeomin 50U/Vial and 100U/Vial*)

- For the treatment of blepharospasm or cervical dystonia (spasmodic torticollis).

INDACATEROL AND GLYCOPYRRONIUM (*Ultibro Breezhaler 110mcg/50mcg Capsule for Inhalation*)

- See [Long-Acting Beta₂-Agonists/Long-Acting Muscarinic Antagonists](#)

INDACATEROL, GLYCOPYRRONIUM AND MOMETASONE FUROATE (*Enerzair Breezhaler 150mcg/50mcg/160mcg Capsule for Inhalation*)

- For the treatment of asthma in adult patients not adequately controlled with a maintenance combination of a long-acting beta₂-agonist (LABA) and a medium or high dose of an inhaled corticosteroid (ICS) who experienced one or more asthma exacerbations in the previous 12 months.

Clinical Note:

- Asthma exacerbation is defined as: worsening signs or symptoms of asthma (shortness of breath, cough, wheezing or chest tightness and progressive decrease in lung function) requiring administration of systemic corticosteroids for at least three days, or asthma-related hospitalization

INDACATEROL, IN COMBINATION (*Ateectura Breezhaler 150mcg/80mcg, 150mcg/160mcg and 150mcg/320mcg Capsule for Inhalation*)

- See [Long-Acting Beta₂-Agonists/Inhaled Corticosteroids](#)

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

INFLIXIMAB (*Avsola, Inflectra and Renflexis 100mg Powder For Injection*)

ANKYLOSING SPONDYLITIS

- For the treatment of patients with moderate to severe ankylosing spondylitis (Bath AS Disease Activity Index (BASDAI) score ≥ 4 on 10 point scale) who:
 - Have axial symptoms¹ and who have failed to respond to the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each, or in whom NSAIDs are contraindicated;
OR
 - Have peripheral symptoms and who have failed to respond to, or have contraindications to, the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each and have had an inadequate response to an optimal dose or maximal tolerated dose of a DMARD.
- Must be prescribed by a rheumatologist or prescriber with a specialty in rheumatology.
- Requests for renewal must include information showing the beneficial effects of the treatment, specifically:
 - A decrease of at least 2 points on the BASDAI scale, compared with the pre-treatment score; OR
 - Patient and expert opinion of an adequate clinical response as indicated by a significant functional improvement (measured by outcomes such as HAQ or "ability to return to work").

Claim Notes:

- Initial period 6 months.
 - Maximum dose 5mg/kg at 0, 2, and 6 weeks then every 6-8 weeks thereafter.
 - Concurrent use of biologics not approved.
1. Patients with recurrent uveitis (2 or more episodes within 12 months) as a complication of axial disease, do not require a trial of 2 NSAIDs.

CROHN'S DISEASE

- For treatment of Crohn's disease in patients with moderate to severe active disease refractory to 5-ASA products AND glucocorticoids (e.g., prednisone) AND immunosuppressive therapy (azathioprine or 6-mercaptopurine or methotrexate)¹.
 - Initial approval of infliximab will be for a single infusion of 5mg/kg/dose. A second infusion may be warranted in patients not responding to the first infusion or in patients responding initially but then worsening before maintenance therapy is effective. Request for approval beyond induction therapy will be considered on a case by case basis.
 - In patients with fistulizing disease who have actively draining perianal or enterocutaneous fistula(e) that have recurred or persisted despite a course of appropriate antibiotic therapy (e.g., metronidazole +/- ciprofloxacin for a minimum of 3 weeks) AND immunosuppressive therapy (azathioprine or 6-mercaptopurine or methotrexate)¹.
 - Initial approval is for three infusions of infliximab of 5mg/kg/dose at 0, 2 and 6 week intervals.
1. Patients who are very ill and not candidates for surgery may qualify for infliximab therapy without a trial of AZA, 6-MP or MTX, as they may require a more rapid onset of response.

Note:

- Requires a written request by a gastroenterologist or physician with a specialty in gastroenterology.

PSORIASIS

- For patients with severe, debilitating chronic plaque psoriasis who meet all of the following:
 - Body surface area (BSA) involvement of $>10\%$ and/or significant involvement of the face, hands, feet or genitals;

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Failure to, contraindication to or intolerant of methotrexate and cyclosporine;
- Failure to, intolerant of or unable to access phototherapy;
- Written request of a dermatologist or prescriber with a specialty in dermatology.
- Continued coverage is dependent on evidence of improvement, specifically:
 - A $\geq 75\%$ reduction in the Psoriasis Area and Severity Index (PASI) score; or
 - A $\geq 50\%$ reduction in PASI with a ≥ 5 -point improvement in DLQI (Dermatology Life Quality Index); or
 - Significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals.

Clinical Note:

- Treatment should be discontinued if a response has not been demonstrated after 12 weeks.

Claim Notes:

- Concurrent use of biologics not approved.
- Initial approval for a maximum of 12 weeks
- Dosage restricted to infliximab 5mg/kg 0, 2 and 6 weeks then every 8 weeks

PSORIATIC ARTHRITIS

- For the treatment of patients with predominantly axial psoriatic arthritis who are refractory, intolerant or have contraindications to the sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each.
- For the treatment of patients with predominantly peripheral psoriatic arthritis who are refractory, intolerant or have contraindications to:
 - The sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each; AND
 - Methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age) for a minimum of 8 weeks; AND
 - Leflunomide for a minimum of 10 weeks or sulfasalazine for a minimum of 3 months.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- 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 to treatments. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial period 3 months.
- Maximum dose 5mg/kg 0, 2 and 6 weeks then every 8 weeks
- Renewal approval: 1 year. Confirmation of continued response required.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

RHEUMATOID ARTHRITIS

- For the treatment of severely active rheumatoid arthritis, in combination with methotrexate or other disease-modifying antirheumatic drugs (DMARDs), in adult patients who are refractory or intolerant to:
 - Methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age), or use in combination with another DMARD, for a minimum of 12 weeks;
- AND
- Methotrexate in combination with at least two other DMARDs, such as hydroxychloroquine and sulfasalazine, for a minimum of 12 weeks.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- Optimal treatment response to DMARDs may take up to 24 weeks, however coverage of a biologic therapy can be considered if no improvement is seen after 12 weeks of triple DMARD use.
- If patient factors (e.g. intolerance) prevent the use of triple DMARD therapy, these must be described and dual therapy with DMARDs must be tried.
- 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. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial Approval: 6 months.
- 3mg/kg/dose at 0, 2 and 6 weeks, then every 8 weeks thereafter.
- Renewal Approval: 1 year. Confirmation of continued response is required.

ULCERATIVE COLITIS

- For the treatment of patients with moderately to severely active ulcerative colitis who have a partial Mayo score > 4 , and a rectal bleeding subscore ≥ 2 and are:
 - refractory or intolerant to conventional therapy (i.e. 5-ASA for a minimum of 4 weeks, and prednisone ≥ 40 mg daily for two weeks or IV equivalent for one week); OR
 - corticosteroid dependent (i.e. cannot be tapered from corticosteroids without disease recurrence; or have relapsed within three months of stopping corticosteroids; or require two or more courses of corticosteroids within one year.)
- Renewal requests must include information demonstrating the beneficial effects of the treatment, specifically:
 - a decrease in the partial Mayo score ≥ 2 from baseline, AND
 - a decrease in the rectal bleeding subscore ≥ 1 .

Clinical Notes:

- 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. The nature of intolerance(s) must be clearly documented.
- Patients with severe disease do not require a trial of 5-ASA

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Notes:

- Must be prescribed by a gastroenterologist or physician with a specialty in gastroenterology.
- Concurrent use of biologics not approved.
- Initial Approval: 16 weeks.
- Renewal Approval: 1 year.

INOTERSEN (Tegsedi 189mg/mL Prefilled Syringe)**POLYNEUROPATHY IN HEREDITARY TRANSTHYRETIN-MEDIATED AMYLOIDOSIS**

- For the treatment of polyneuropathy in adult patients with hereditary transthyretin-mediated amyloidosis (hATTR) who meet all of the following criteria:
 - Confirmed genetic diagnosis of hATTR
 - Symptomatic with early-stage neuropathy
 - Does not have New York Heart Association class III or IV heart failure
 - Has not previously undergone a liver transplant

Discontinuation Criteria:

- The patient is permanently bedridden and dependent on assistance for basic activities of daily living
OR
- The patient is receiving end-of-life care.

Clinical Note:

- Symptomatic early-stage neuropathy is defined as polyneuropathy disability stage I to IIIB or familial amyloidotic polyneuropathy stage I or II.

Claim Notes:

- The patient must be under the care of a physician with experience in the diagnosis and management of hATTR.
- Combination therapy with other interfering ribonucleic acid drugs or transthyretin stabilizers used to treat hATTR will not be reimbursed.
- Initial Approval: 9 months.
- Renewal Approval: 12 months. Confirmation of continued response is required.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904551
 - 00904552
 - 00904553

INSULIN DETEMIR (Levemir 100iu/mL Penfill and FlexTouch Prefilled Pen)

- For the treatment of patients who have been diagnosed with Type 1 or Type 2 diabetes requiring insulin and have previously taken NPH and/or premix insulin daily at optimal dosing

AND

- have experienced unexplained nocturnal hypoglycemia at least once a month despite optimal management

OR

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- have documented severe or continuing systemic or local allergic reaction to existing insulin(s)
- For the treatment of pediatric and adolescent patients (under 18 years of age) with Type 1 diabetes.
- For the treatment of pregnant individuals with Type 1 or Type 2 diabetes requiring insulin.

INTERFERON BETA-1A (*Avonex PS 30mcg/0.5mL Injection and Rebif 22mcg Multidose Cartridges, 22mcg/0.5mL Injection, 44mcg Multidose Cartridges and 44mcg/0.5mL Injection*)
INTERFERON BETA-1B (*Betaseron 0.3mg/Vial Injection*)

Prescribed by a neurologist with experience in the treatment of multiple sclerosis for patients who meet the following criteria:

Treatment initiation:

- Diagnosis of multiple sclerosis with a relapsing course*.
 - Includes relapsing-remitting MS and secondary progressive MS with clear superimposed relapses;
 - Does not include primary progressive MS, progressive- relapsing or secondary progressive MS without relapses;
 - and
 - Disability judged to be equivalent to Expanded Disability Status Score (EDSS) of 5.5 or less.

Renewal:

- EDSS not greater than 6.0 for at least 12 months in the absence of relapses.
- Patients must be assessed for compliance and for any therapy related side effects that are intolerable.

Exclusions:

- Concurrent illness likely to alter compliance or substantially reduce life expectancy.
- Planned pregnancy, pregnancy or breast-feeding.
- Active and severe depression.

* Relapsing course is defined as evidence of one relapse in the past 18 months or two relapses in the past 3 years.

***IPRATROPIUM BROMIDE** (*Atrovent 125mcg/mL and 250mcg/mL Inhaled Solutions and generic brands*)

- See [Wet Nebulization Solutions](#)

***IPRATROPIUM BROMIDE, IN COMBINATION** (*Combivent Inhaled Solution and generic brands*)

- See [Wet Nebulization Solutions](#)

***ISAVUCONAZOLE** (*Cresemba 100mg Capsule and 200mg Vial*)

- For the treatment of adult patients with invasive aspergillosis who have a contraindication, intolerance or have failed to respond to oral voriconazole and caspofungin.
- For the treatment of adult patients with invasive mucormycosis.

Claim Notes:

- Must be prescribed by a hematologist or specialist in infectious diseases or medical microbiology.
- Initial requests will be approved for a maximum of 3 months.
- Claims for Cresemba 200mg Vial that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00904516

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

***ITRACONAZOLE** (*Sporanox 10mg/mL Oral Solution and generic brands*)

- For the treatment of immunocompromised adult patients with oral and/or esophageal candidiasis.

Clinical Note:

- Itraconazole oral solution is not interchangeable with itraconazole capsules due to differences in bioavailability

IVABRADINE HYDROCHLORIDE (*Lancora 5mg and 7.5mg Tablet*)

- For the treatment of adult patients with New York Heart Association (NYHA) classes II or III stable chronic heart failure to reduce the incidence of cardiovascular death and hospitalization, administered in combination with standard chronic heart failure therapies, who meet all of the following criteria:
 - reduced left ventricular ejection fraction (LVEF) (<35%)
 - sinus rhythm with a resting heart rate ≥ 77 beats per minute (bpm)
 - at least one hospitalization due to heart failure in the past year
 - NYHA class II to III symptoms despite at least four weeks of optimal treatment of the following:
 - a stable dose of an angiotensin converting enzyme inhibitor (ACEI) or an angiotensin II receptor blocker (ARB); and
 - a stable dose of a beta blocker; and
 - an aldosterone antagonist

Clinical Notes:

- Resting heart rate must be documented as ≥ 77 bpm on average using either an ECG on at least three separate visits or by continuous monitoring.
- For patients who have not received four weeks of therapy with an ACEI/ARB, beta blocker or aldosterone antagonist due to an intolerance or contraindication, details must be provided.

Claim Note:

- Patients should be under the care of a specialist experienced in the treatment of heart failure for patient selection, titration, follow-up and monitoring.

IXEKIZUMAB (*Taltz 80mg/mL Autoinjector and Prefilled Syringe*)

PSORIASIS

- For patients with severe, debilitating chronic plaque psoriasis who meet all of the following:
 - Body surface area (BSA) involvement of $>10\%$ and/or significant involvement of the face, hands, feet or genitals;
 - Failure to, contraindication to or intolerant of methotrexate and cyclosporine;
 - Failure to, intolerant of or unable to access phototherapy;
 - Written request of a dermatologist or prescriber with a specialty in dermatology.
- Continued coverage is dependent on evidence of improvement, specifically:
 - A $\geq 75\%$ reduction in the Psoriasis Area and Severity Index (PASI) score; or
 - A $\geq 50\%$ reduction in PASI with a ≥ 5 -point improvement in DLQI (Dermatology Life Quality Index); or
 - Significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Note:

- Treatment should be discontinued if a response has not been demonstrated after 12 weeks.

Claim Notes:

- Concurrent use of biologics not approved.
- Initial approval for a maximum of 12 weeks.
- Approvals will be for 160 mg at week 0, followed by 80 mg at weeks 2, 4, 6, 8, 10, and 12 then 80 mg every four weeks.
- Renewal approval: 1 year.

PSORIATIC ARTHRITIS

- For the treatment of patients with predominantly axial psoriatic arthritis who are refractory, intolerant or have contraindications to the sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each.
- For the treatment of patients with predominantly peripheral psoriatic arthritis who are refractory, intolerant or have contraindications to:
 - The sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each; AND
 - Methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age) for a minimum of 8 weeks; AND
 - Leflunomide for a minimum of 10 weeks or sulfasalazine for a minimum of 3 months.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- 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 to treatments. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.
- Initial approval for a maximum of 12 weeks.
- Approvals will be for 160mg at week 0, followed by 80mg every 4 weeks.
- Renewal approval: 1 year. Confirmation of continued response required.

***LACTULOSE (667mg/mL Oral Liquid, generic brands)**

- For portal systemic encephalopathy.
- For pneumatosis cystoides intestinalis.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

LANADELUMAB (*Takhzyro 300mg/2mL Vial and 300mg/2mL Prefilled Syringe*)

- For the routine prevention of attacks of type I or II hereditary angioedema (HAE) in patients 12 years of age and older who have experienced at least three HAE attacks within any four-week period and required the use of an acute injectable treatment.

Discontinuation Criteria:

- No reduction in the number of HAE attacks for which acute injectable treatment was received during the first three months of treatment with lanadelumab compared to the number of attacks observed before initiating treatment with lanadelumab;

OR

- Increase in the number of HAE attacks for which acute injectable treatment was received compared to the number of attacks before initiating treatment with lanadelumab.

Clinical Note:

- The pre-treatment attack rate must be provided for those patients who are already receiving long-term prophylactic treatment for HAE and intend to transition to lanadelumab.

Claim Notes:

- Must be prescribed by a physician experienced in the diagnosis and treatment of HAE.
- Combination use of Takhzyro (lanadelumab) with other long-term prophylactic treatment of HAE (e.g., C1 esterase inhibitor) will not be funded.
- Approvals will be for a maximum of 300 mg every two weeks.
- Initial approval period: 3 months.
- Renewal approval period: 6 months.
- Claims for Takhzyro 300mg/2mL vial and 300mg/2mL prefilled syringe that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - Takhzyro 300mg/2mL Vial
 - 00904577
 - 00904578
 - Takhzyro 300mg/2mL Prefilled Syringe
 - 00904638
 - 00904639

***LANSOPRAZOLE** (*Prevacid FasTab 15mg, 30mg Tablet*)

- For patients who require the use of a proton pump inhibitor and require administration through a feeding tube. **[Criteria code 37]**
- For patients 19 years of age and younger, who require the use of a proton pump inhibitor and who cannot use a tablet or capsule. **[Criteria code 38]**

***LANSOPRAZOLE** (*Prevacid 15mg, 30mg Capsule and generic brands*)

- See [Proton Pump Inhibitors](#)

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

*LAPATINIB (Tykerb 250mg Tablet)

- For the treatment of patients with unresectable locally advanced or metastatic HER2-positive breast cancer in combination with capecitabine for use as:
 - First line therapy following disease relapse during or within six months of completing adjuvant treatment with trastuzumab or trastuzumab emtansine; or
 - Second line therapy following disease progression during treatment with trastuzumab, with or without pertuzumab, in the advanced setting.

Clinical Note:

- Patients should have a good performance status.

*LAROTRECTINIB (Vitrakvi 25mg and 100mg Capsule and 20mg/mL Oral Liquid)

LOCALLY ADVANCED UNRESECTABLE OR METASTATIC SOLID TUMORS WITH A NEUROTROPHIC TYROSINE RECEPTOR KINASE (NTRK) GENE FUSION

- For the treatment of adult and pediatric patients with locally advanced unresectable or metastatic solid tumors with NTRK gene fusion without a known acquired resistance mutation. Patient is not a candidate for surgery and/or radiation due to risk of substantial morbidity and have no satisfactory treatment options.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.
- Brain metastases are stable, if present.
- Patients with prior progression on an NTRK inhibitor are not eligible.

Claim Note:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - Vitrakvi 100mg Cap - 00900013
 - Vitrakvi 20mg/mL O/L - 00900014

LEDIPASVIR AND SOFOSBUVIR (Harvoni 90mg/400mg Tablet)

- For treatment-naïve or treatment-experienced adult patients with chronic hepatitis C virus (HCV) who meet the following criteria:

Approval Period and Regimen

Genotype 1

- Treatment-naïve without cirrhosis, who have pre-treatment HCV RNA level < 6 million IU/mL and mono-HCV infected only

- 8 weeks

Genotype 1

- Treatment-naïve without cirrhosis, who have pre-treatment HCV RNA level ≥ 6 million IU/mL
- Treatment-naïve with compensated cirrhosis
- Treatment-naïve with advanced liver fibrosis (Fibrosis stage F3-F4)
- Treatment-experienced without cirrhosis

- 12 weeks

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Approval Period and Regimen

- HCV/HIV co-infected without cirrhosis or with compensated cirrhosis

Genotype 1

- Treatment-experienced with compensated cirrhosis
- 24 weeks

Genotype 1

- Decompensated cirrhosis
- 12 weeks in combination with ribavirin
- Liver transplant recipients without cirrhosis or with compensated cirrhosis
- Patients must also meet all of the following criteria:
 - Must be prescribed by a hepatologist, gastroenterologist, or infectious disease specialist (or other physician experienced in treating a patient with hepatitis C infection);
 - Lab-confirmed hepatitis C genotype 1;
 - Quantitative HCV RNA value within the last 6 months;
 - Fibrosis stage must be provided.

Clinical Notes:

1. Treatment-experienced is defined as a patient who has been previously treated with a peginterferon/ribavirin regimen, including regimens containing HCV protease inhibitors and who has not experienced an adequate response.
2. Acceptable methods for the measurement of fibrosis score include Fibrotest, liver biopsy, transient elastography (FibroScan®), serum biomarker panels (such as AST-to-Platelet Ratio Index or Fibrosis-4 score) either alone or in combination.
3. Compensated cirrhosis is defined as a CTP score of 5 to 6 (Class A) and decompensated cirrhosis as a CTP score of 7 or above (Class B or C).
4. Re-treatment for direct-acting antiviral failures will be considered on a case-by-case basis.

Claim Note:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the following PINs:
 - 00904032
 - 00904033
- Claims will be limited to a 28-day supply.

** [Criteria Code 34] has been added for use effective December 1, 2019 to allow payment of a patient's initial 28 day supply only. Criteria code 34 should be provided by the prescribing physician only.

***LENALIDOMIDE** (*Revlimid 2.5mg, 5mg, 10mg, 15mg, 20mg, 25mg Capsule and generic brands*)

MYELOYDYSPLASTIC SYNDROME (MDS)

- As a single agent in adult myelodysplastic syndrome (MDS) patients with transfusion dependent anemia due to low or intermediate-1 risk MDS associated with a deletion 5q cytogenetic abnormality with or without additional cytogenetic abnormalities.

MULTIPLE MYELOMA (MM-AOPT)

- For the treatment of relapsed or refractory multiple myeloma when used:
 - In combination with dexamethasone for patients who have received at least one prior treatment; or

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- In combination with carfilzomib and dexamethasone (KRd regimen) for patients who have received at least one prior treatment; or
- In combination with daratumumab and dexamethasone (DRd regimen) for patients who have received at least one prior treatment.

NEWLY DIAGNOSED MULTIPLE MYELOMA POST-AUTOLOGOUS STEM CELL TRANSPLANT (NDMM POST-ASCT)

- For the maintenance treatment of patients with newly diagnosed multiple myeloma who have stable or improved disease following autologous stem-cell transplantation (ASCT) and no evidence of disease progression.

MULTIPLE MYELOMA NOT ELIGIBLE FOR AUTOLOGOUS STEM CELL TRANSPLANT (MM-TNE)

- As first-line treatment for newly diagnosed patients with multiple myeloma who are not eligible for autologous stem cell transplantation when used:
 - In combination with dexamethasone, with or without bortezomib; or
 - In combination with daratumumab and dexamethasone

MULTIPLE MYELOMA PRIOR TO AUTOLOGOUS STEM CELL TRANSPLANT (MM PRE-ASCT)

- For the treatment of patients with multiple myeloma when used in combination with bortezomib and dexamethasone as induction therapy prior to autologous stem cell transplant.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be continued until unacceptable toxicity or disease progression.

***LENVATINIB (*Lenvima 4mg, 8mg, 10mg, 12mg, 14mg, 20mg, 24mg Compliance Pack*)**

ADVANCED AND METASTATIC RENAL CELL CARCINOMA

- In combination with pembrolizumab for the treatment of adult patients with advanced (not amenable to curative surgery or radiation) or metastatic renal cell carcinoma who have not had prior systemic therapy for metastatic disease.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should continue until disease progression or unacceptable toxicity (can be continued as monotherapy after completing 2 years of combination therapy with pembrolizumab).
- If pembrolizumab or lenvatinib is discontinued for toxicity, the other agent can be continued at the discretion of the physician.
- Patients are eligible for one of pembrolizumab with lenvatinib or pembrolizumab with axitinib in this setting. If intolerant to one tyrosine kinase inhibitor (TKI), patient may be switched to an alternate TKI, provided there has been no progression.
- Patients who received pembrolizumab in the adjuvant setting are eligible for treatment provided there was a disease-free interval of at least six months.
- If patient requires and qualifies for re-treatment with pembrolizumab, lenvatinib may also be given at the discretion of the treating physician.

ADVANCED ENDOMETRIAL CARCINOMA

- In combination with pembrolizumab for the treatment of adult patients with advanced endometrial carcinoma that is not microsatellite instability high (MSI-H) or mismatch repair deficient (dMMR), who have disease progression following prior platinum-based systemic therapy and are not candidates for curative surgery or radiation.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

- Patients should have a good performance status.
- Treatment should continue until disease progression or unacceptable toxicity (can be continued as monotherapy after completing 2 years of combination therapy with pembrolizumab).
- Confirmation that patient does not have MSI-H or dMMR disease must be done prior to initiating treatment.
- No active CNS metastases (eligible if treated/stable).
- If pembrolizumab or lenvatinib is discontinued for toxicity, the other agent can be continued at the discretion of the physician.
- If patient requires and qualifies for re-treatment with pembrolizumab, lenvatinib can also be given at the discretion of the treating physician.

LOCALLY RECURRENT OR METASTATIC, PROGRESSIVE, RADIOACTIVE-IODINE-REFRACTORY DIFFERENTIATED THYROID CANCER (DTC) (LENVIMA 10MG,14MG, 20MG AND 24MG COMPLIANCE PACK)

- For the treatment of patients with locally recurrent or metastatic, progressive, radioactive-iodine-refractory differentiated thyroid cancer (DTC). Treatment should be for patients with good performance status and who otherwise meet the eligibility criteria of the SELECT trial and should continue until treatment progression or unacceptable toxicity.

UNRESECTABLE OR METASTATIC HEPATOCELLULAR CARCINOMA (LENVIMA 4MG, 8MG AND 12MG COMPLIANCE PACK)

- For the treatment of adult patients with unresectable or metastatic hepatocellular carcinoma as either first-line treatment, or second-line treatment following atezolizumab in combination with bevacizumab, who meet all the following criteria:
 - Child-Pugh class status of A
 - ECOG performance status of 0 or 1
 - Less than 50% liver involvement and no invasion of the bile duct or main portal vein
 - No brain metastases or prior liver transplantation

Clinical Notes:

- Treatment should be continued until disease progression or unacceptable toxicity.
- Patients who are unable to tolerate lenvatinib may be switched to sorafenib if there is no disease progression and provided all other funding criteria are met.
- Patients with disease progression on lenvatinib are not eligible for reimbursement of sorafenib.

LETERMOVIR (*Prevymis 240mg and 480mg Tablet and 240mg and 480mg IV Solution*)

- For the prevention of cytomegalovirus (CMV) infection in adult CMV-seropositive recipients [R+] of an allogeneic hematopoietic stem cell transplant (HSCT) who have undetectable CMV viremia at baseline and meet one of the following criteria:
 - umbilical cord blood as a stem cell source
 - recipient of a haploidentical transplant
 - recipient of T-cell depleted transplant
 - treated with antithymocyte globulin (ATG) for conditioning
 - requiring high-dose steroids or other immunosuppression for acute graft versus host disease (GVHD)
 - treated with ATG for steroid-refractory acute GVHD

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- documented history of CMV disease prior to transplantation

Clinical Note:

- High-dose steroids is defined as the use of greater than or equal to 1 mg/kg/day of prednisone or equivalent dose of another corticosteroid.

Claim Notes:

- Must be prescribed by a medical oncologist, hematologist, or infectious disease specialist or other physician with experience in the management of HSCT.
- Approvals will be for a maximum dose of 480 mg per day.
- Approval period: 100 days per HSCT.

LEVOCARNITINE (*Carnitor 100mg/mL Oral Liquid, generic brands and 330mg Tablet*)

- For the treatment of patients with primary systemic carnitine deficiency.
- For the treatment of patients with an inborn error of metabolism that results in secondary carnitine deficiency.

LEVODOPA/CARBIDOPA (*Duodopa 20mg/5mg Intestinal Gel Cassettes*)

- For the treatment of patients with advanced levodopa-responsive Parkinson's Disease (PD) who meet all of the following criteria:
 - Experiences severe disability with at least 25% of the waking day in the off state and/or ongoing levodopa-induced dyskinesias, despite having tried frequent dosing of levodopa (at least five doses per day).
 - Have received an adequate trial of maximally tolerated doses of levodopa, with demonstrated clinical response.
 - Have failed an adequate trial of the following adjunctive medications, if not contraindicated and/or contrary to the clinical judgment of prescriber: entacapone, a dopamine agonist, a monoamine oxidase-B (MAO-B) inhibitor and amantadine.
 - Must be able to administer the medication and care for the administration port and infusion pump. Alternatively, trained personnel or a care partner must be available to perform these tasks reliably.

Exclusion Criteria:

- Patients with a contraindication to the insertion of a PEG-J tube.
- Patients with severe psychosis or dementia.

Renewal Criteria:

- Patients continue to demonstrate a significant reduction in the time spent in the off state and/or in ongoing levodopa-induced dyskinesias, along with and an improvement in the related disability.

Clinical Note:

- Time in the off state, frequency of motor fluctuations, and severity of associated disability should be assessed by a movement disorder subspecialist and be based on an adequate and reliable account from longitudinal specialist care, clinical interview of a patient and/or care partner, or motor symptom diary.

Claim Notes:

- Must be prescribed by a movement disorder subspecialist who has appropriate training in the use of Duodopa and is practicing in a movement disorder clinic that provides ongoing management and support for patients

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

receiving treatment with Duodopa.

- Approval period: 1 year.

LEVODOPA AND CARBIDOPA AND ENTACAPONE (*Stalevo 50mg, 75mg, 100mg, 125mg, 150mg Tablet*)

- For the treatment of Parkinson's disease as adjunctive therapy in patients who:
 - are not well controlled and are experiencing significant "wearing off" symptoms despite optimal therapy with levodopa/carbidopa;
 - were not well controlled and experienced significant "wearing off" symptoms despite optimal therapy with levodopa/carbidopa and are currently using levodopa/carbidopa and entacapone separately.

***LEVOFLOXACIN** (*Levaquin 250mg, 500mg Tablet and generic brands*)

- See [Fluoroquinolones, Respiratory](#)

LINEZOLID (*Zyvoxam 600mg Tablet and generic brands*)

- Written request from an infectious disease specialist or prescriber with a specialty in infectious diseases.
- For the treatment of proven vancomycin-resistant enterococci (VRE) infections.
- For the treatment of proven methicillin-resistant staphylococcus aureus or epidermidis (MRSA/MRSE) infections in those patients who are unresponsive to, or intolerant of vancomycin.

LINAGLIPTIN (*Trajenta 5mg Tablet*)

- For the treatment of Type II diabetes for patients with:
 - inadequate glycemic control on metformin and a sulfonylurea; and
 - for whom insulin is not an option.

LINAGLIPTIN/METFORMIN (*Jentaduetto 2.5mg/500mg, 2.5mg/850mg and 2.5mg/1000mg Tablet*)

- For the treatment of Type II diabetes for patients:
 - who are already stabilized on therapy with metformin, a sulfonylurea and linagliptin to replace the individual components of linagliptin and metformin; and
 - for whom insulin is not an option.

LISDEXAMFETAMINE (*Vyvanse 10mg, 20mg, 30mg, 40mg, 50mg, and 60mg Capsule and 10mg, 20mg, 30mg, 40mg, 50mg, and 60mg Chewable Tablet*)

- For treatment of patients with attention deficit hyperactivity disorder who have tried extended-release methylphenidate, dexamphetamine or mixed salts amphetamine with unsatisfactory results.

Claim Note:

- The maximum dose reimbursed is 60 mg daily.

LONG-ACTING BETA₂-AGONISTS (*Formoterol, Salmeterol*)

ASTHMA

- For the treatment of moderate to severe asthma in patients who:
 - are compliant with inhaled corticosteroids at optimal doses; and
 - require additional symptom control, (e.g., cough, awakening at night, missing activities such as school, work or social activities because of asthma symptoms); and
 - require increasing amounts of short-acting beta₂-agonists, indicative of poor control.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Products and Strengths Approved:

Oxeze • 6mcg/Dose, 12mcg/Dose Turbuhaler

Serevent • 50mcg/dose Diskus

CHRONIC OBSTRUCTIVE PULMONARY DISEASE

- For the treatment of chronic obstructive pulmonary disease (COPD), as defined by spirometry, in patients who experience:
 - persistent symptoms, as defined by Medical Research Council (MRC) Dyspnea Scale of at least Grade 3 or a COPD Assessment test (CAT) score of at least 10 and have a post-bronchodilator FEV₁ less than 80% predicted; OR
 - two or more moderate exacerbations of COPD in the previous year requiring treatment with antibiotics and/or systemic corticosteroids; OR
 - at least one acute severe exacerbation of COPD requiring hospitalization.

Clinical Note:

- COPD is defined by spirometry as a post-bronchodilator FEV₁/FVC ratio less than 0.7. Spirometry reports from any point in time will be accepted. If spirometry cannot be obtained, reasons must be clearly explained and other evidence of COPD severity provided (i.e. MRC Dyspnea Scale grade).

Claim Note:

- Requests for combination therapy of single agent long-acting bronchodilators, i.e. long-acting beta-2 agonist (LABA) and long-acting muscarinic antagonist (LAMA), will not be considered. Products which combine a LABA/LAMA in a single device are available as special authorization benefits with their own criteria.

Products and Strengths Approved:

Serevent • 50mcg/dose Diskus

LONG-ACTING BETA₂-AGONISTS/INHALED CORTICOSTEROIDS (*Formoterol, In Combination; Indacaterol, in Combination, Salmeterol, In Combination*)

ASTHMA

- For the treatment of moderate to severe asthma in patients who:
 - are compliant with inhaled corticosteroids at optimal doses; and
 - require additional symptom control, (e.g., cough, awakening at night, missing activities such as school, work or social activities because of asthma symptoms); and
 - require increasing amounts of short-acting beta₂-agonists, indicative of poor control.

Products and Strengths Approved:

Ateectura • 150mcg/80mcg, 150mcg/160mcg and 150mcg/320mcg Capsule for Inhalation

Advair and generic brands • 50/100mcg, 50/250mcg and 50/500mcg Diskus
• HFA 25/125 mcg/dose
• HFA 25/250 mcg/dose Inhaler

Breo Ellipta • 100mcg/25mcg and 200mcg/25mcg dry powder for inhalation

Symbicort • 100/6mcg Turbuhaler

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Products and Strengths Approved:

- 200/6mcg Turbuhaler

Zenhale

- 5/100mcg and 5/200mcg

CHRONIC OBSTRUCTIVE PULMONARY DISEASE

- For the treatment of chronic obstructive pulmonary disease (COPD), as defined by spirometry, in combination with a long-acting muscarinic antagonist (LAMA), in patients who experience inadequate control while being treated with a long-acting beta-2 agonist/long-acting muscarinic antagonist (LABA/LAMA).
- For the treatment of patients with asthma / chronic obstructive pulmonary disease (ACO) overlap, based on patient history and lung function studies indicating an ACO diagnosis.
 - Please provide details to support the ACO diagnosis (patient symptoms, risk factors, spirometry etc.).

Clinical Notes:

- COPD is defined by spirometry as a post-bronchodilator FEV₁/FVC ratio less than 0.7. Spirometry reports from any point in time will be accepted. If spirometry cannot be obtained, reasons must be clearly explained and other evidence of COPD severity provided (i.e. MRC Dyspnea Scale grade).
- Inadequate control while being treated with a LABA/LAMA for at least two months is defined as persistent symptoms, or experiencing two or more exacerbations of COPD in the previous year requiring treatment with antibiotics and/or systemic corticosteroids or at least one exacerbation of COPD requiring hospitalization.

Products and Strengths Approved:

Advair and generic brands

- 50/100mcg Diskus
- 50/250mcg Diskus
- 50/500mcg Diskus

Breo Ellipta

- 100mcg/25mcg dry powder for inhalation

Symbicort

- 100/6mcg Turbuhaler
- 200/6mcg Turbuhaler

LONG-ACTING BETA₂-AGONISTS/LONG-ACTING MUSCARINIC ANTAGONISTS (*Formoterol and Acclidinium Bromide, Indacaterol and Glycopyrronium Bromide, Olodaterol and Tiotropium Bromide and Vilanterol and Umeclidinium Bromide*)

- For the treatment of chronic obstructive pulmonary disease (COPD), as defined by spirometry, in patients who experience inadequate control while being treated with either a long-acting beta-2 agonist (LABA) or long-acting muscarinic antagonist (LAMA).

Clinical Notes:

- COPD is defined by spirometry as a post-bronchodilator FEV₁/FVC ratio less than 0.70. Spirometry reports from any point in time will be accepted. If spirometry cannot be obtained, reasons must be clearly explained, and other evidence of COPD severity provided (i.e. Medical Research Council (MRC) Dyspnea Scale grade).
- Inadequate control is defined as persistent symptoms (e.g. MRC Dyspnea Scale of at least grade 3 or COPD Assessment test (CAT) score of at least 10) after at least one month of a LAMA or LABA.
- LABA/LAMA combinations are not intended to be used with an inhaled corticosteroid (ICS) unless criteria for triple inhaled therapy (LABA/LAMA/ICS) is met.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

LONG-ACTING MUSCARINIC ANTAGONISTS (*Aclidinium Bromide, Glycopyrronium Bromide, Tiotropium Bromide, Tiotropium Bromide Monohydrate and Umeclidinium*)

- For the treatment of chronic obstructive pulmonary disease (COPD), as defined by spirometry, in patients who experience:
 - persistent symptoms, as defined by Medical Research Council (MRC) Dyspnea Scale of at least Grade 3 or a COPD Assessment test (CAT) score of at least 10 and have a post-bronchodilator FEV₁ less than 80% predicted; OR
 - two or more moderate exacerbations of COPD in the previous year requiring treatment with antibiotics and/or systemic corticosteroids; OR
 - at least one acute severe exacerbation of COPD requiring hospitalization.
- For the treatment of COPD, as defined by spirometry, in combination with a long-acting beta-2 agonist/inhaled corticosteroid (LABA/ICS), for patients who experience inadequate control while being treated with a LABA/ICS or a long-acting beta-2 agonist/long-acting muscarinic receptor antagonists (LABA/LAMA).

Clinical Notes:

- COPD is defined by spirometry as a post-bronchodilator FEV₁/FVC ratio less than 0.7. Spirometry reports from any point in time will be accepted. If spirometry cannot be obtained, reasons must be clearly explained and other evidence of COPD severity provided (i.e. MRC Dyspnea Scale grade).
- Inadequate control while being treated with a LABA/LAMA or LABA/ICS for at least two months is defined as persistent symptoms or experiencing two or more exacerbations of COPD in the previous year requiring treatment with antibiotics and/or systemic corticosteroids or at least one exacerbation of COPD requiring hospitalization.

Claim Note:

- Requests for combination therapy of single agent long-acting bronchodilators, i.e. LABA and LAMA, will not be considered. Products which combine a LABA/LAMA in a single device are available as special authorization benefits with their own criteria.

LORATADINE (*Claritin 10mg Tablet and generic brands*)

- For chronic urticaria, defined as the presence of hives or lesions for longer than six weeks, which has responded to treatment with loratadine.

***LORLATINIB** (*Lorbrena 25mg and 100mg Tablet*)

- As monotherapy for the first-line treatment of adult patients with anaplastic lymphoma kinase (ALK)-positive locally advanced (not amenable to curative therapy) or metastatic non-small cell lung cancer (NSCLC).

Clinical Notes:

- Patients should have a good performance status.
- Treatment should continue until disease progression or unacceptable toxicity.
- Patients must not have had any prior systemic treatment for advanced or metastatic disease.
- Patients are not eligible for subsequent ALK inhibitor therapy following disease progression on lorlatinib.
- Patients may be switched to an alternate ALK inhibitor in the case of intolerance without disease progression.

Claim Note:

- Claims that exceed \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - Lorbrena 100mg Tab

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- 00900025

LUSPATERCEPT (*Reblozyl 25mg and 75mg Vial*)

BETA-THALASSEMIA ANEMIA

- For the treatment of adult patients with RBC transfusion-dependent anemia associated with beta-thalassemia. Patients must be receiving regular transfusions, defined as:
 - 6 to 20 RBC units in the 24 weeks prior to initiating treatment with luspatercept, AND
 - No transfusion-free period greater than 35 days in the 24 weeks prior to initiating treatment with luspatercept.

Renewal Criteria:

- Patients must demonstrate an initial response, defined as a $\geq 33\%$ reduction in transfusion burden (RBC units/time) compared to the pre-treatment baseline RBC transfusion burden, measured over 24 weeks prior to initiating treatment with luspatercept.
- For continued coverage, patients should maintain a reduction in transfusion burden of $\geq 33\%$ compared to the pre-luspatercept transfusion burden.
- Luspatercept should be discontinued if a patient does not respond after nine weeks of treatment (three doses) at the maximum dose.

Claim Notes:

- The patient should be under the care of a specialist with experience in managing patients with beta-thalassemia.
- The maximum dose of luspatercept should not exceed 1.25mg/kg (or 120mg total dose) once every three weeks.
- Initial Approval: 6 months
- Renewal Approval: 1 year
- Claims that exceed \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - Reblozyl 25mg Vial
 - 00904728
 - Reblozyl 75mg Vial
 - 00904729

MYELOYDYSPLASTIC SYNDROMES

- For the treatment of adult patients with red blood cell (RBC) transfusion-dependent anemia associated with very low- to intermediate-risk MDS who have ring sideroblasts and who have failed or are not suitable for erythropoietin-based therapy.

Renewal Criteria:

- Patients should be RBC transfusion independent over a minimum of 16 consecutive weeks within the first 24 weeks of treatment initiation.
- For continued coverage, patients should be RBC transfusion independent over a minimum of 16 consecutive weeks within the previous approval period.

Claims Notes:

- Treatment should be initiated by a specialist with expertise in managing and treating patients with MDS.
- The maximum dose of luspatercept should not exceed 1.75mg/kg (or 168mg total dose) once every three weeks.
- Approval: 6 months

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Claims that exceed \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - See above.

MACITENTAN (*Opsumit 10mg Tablet*)

- For the treatment of patients with Group 1 pulmonary arterial hypertension (PAH) with a World Health Organization (WHO) functional class of at least II.

Clinical Note:

- The diagnosis of PAH should be confirmed by right heart catheterization.

Claim Notes:

- Must be prescribed by, or in consultation with, a physician experienced in the treatment of PAH.
- Combined use of more than one endothelin receptor antagonists will not be reimbursed.
- The maximum dose of macitentan that will be reimbursed is 10mg daily.

MAGNESIUM GLUCOHEPTONATE (*5mg/mL Solution and generic brands*)

- For the treatment of hypomagnesemia.

MECASERMIN (*Increlex 10mg/mL Vial*)

- For the treatment of growth failure in children and adolescents from 2 to 18 years with confirmed severe primary insulin-like growth factor-1 deficiency (SPIGFD) who meet the following criteria:
 - Epiphyseal closure has not yet occurred; AND
 - Have a confirmed diagnosis of SPIGFD, defined by:
 - a known genetic mutation recognized as a cause of SPIGFD, AND/OR
 - has clinical and biochemical features of SPIGFD

Renewal Criteria:

- Treatment with mecasermin must be discontinued upon the occurrence of any of the following:
 - Height velocity is less than 1cm per 6 months or less than 2cm per year, OR
 - Bone age is more than 16 years in boys and 14 years in girls.

Claim Notes:

- The patient must be under the care of a pediatric endocrinologist
- Mecasermin must not be prescribed concomitantly with recombinant GH treatment
- Approvals: 1 year
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00900015
 - 00900016

MEPOLIZUMAB (*Nucala 100mg/mL Powder for Injection, 100mg/mL Autoinjector and 100mg/mL Pre-filled Syringe*)

- For the adjunctive treatment of severe eosinophilic asthma in adult patients who are inadequately controlled with high dose inhaled corticosteroids and one or more additional asthma controller(s) (e.g., long-acting beta-agonist),

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

and meets one of the following criteria:

- blood eosinophil count of $\geq 0.3 \times 10^9/L$ within the past 12 months and has experienced two or more clinically significant asthma exacerbations in the past 12 months, OR
- blood eosinophil count of $\geq 0.15 \times 10^9/L$ and is receiving maintenance treatment with oral corticosteroids (OCS).

Initial Discontinuation Criteria:

- Baseline asthma control questionnaire score has not improved at 12 months since the initiation of treatment, OR
- No decrease in the daily maintenance OCS dose in the first 12 months of treatment, OR
- Number of clinically significant asthma exacerbations has increased within the previous 12 months.

Subsequent Discontinuation Criteria:

- Baseline asthma control questionnaire score achieved after the first 12 months of therapy has not been maintained subsequently, OR
- Reduction in the daily maintenance OCS dose achieved after the first 12 months of treatment is not maintained subsequently, OR
- Number of clinically significant asthma exacerbations has increased within the previous 12 months.

Clinical Notes:

1. A baseline and annual assessment of asthma symptom control using a validated asthma control questionnaire must be provided.
2. High-dose inhaled corticosteroids is defined as greater than or equal to 500 mcg of fluticasone propionate or equivalent daily dose.
3. A clinically significant asthma exacerbation is defined as worsening of asthma such that the treating physician elected to administer systemic glucocorticoids for at least 3 days or the patient visited an emergency department or was hospitalized.

Claim Notes:

- Must be prescribed by a respirologist, clinical immunologist, allergist or internist with experience in treating severe eosinophilic asthma.
- Combined use of mepolizumab with other biologics used to treat asthma will not be reimbursed.
- Approvals will be for a maximum of 100 mg every four weeks.
- Initial approval period: 1 year.
- Renewal approval period: 1 year.

***METHADONE (*Metadol 1mg, 5mg, 10mg, 25mg Tablet*)**

- For the management of severe chronic or malignant pain as an alternative to other opiates.
- Written request of a physician authorized to prescribe methadone.

Clinical Note:

- In the case of comorbid opioid use disorder (past or current), methadone oral liquid would normally be prescribed as per treatment standards. If methadone tablets are requested in this context, a specialist consult may be required.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

METHYLPHENIDATE (*Biphentin 10mg, 15mg, 20mg 30mg, 40mg, 50mg, 60mg and 80mg Capsule and generic brands*)

- For the treatment of patients with attention deficit hyperactivity disorder who have tried other forms of extended-release methylphenidate with unsatisfactory results.

Claim Note:

- The maximum dose reimbursed is 80mg daily.

METHYLPHENIDATE HYDROCHLORIDE (*Foquest 25mg, 35mg, 45mg, 55mg, 70mg, 85mg and 100mg Capsule*)

- For the treatment of patients with attention deficit hyperactivity disorder (ADHD) who have tried other forms of extended-release methylphenidate with unsatisfactory results.

Claim Note:

- The maximum dose reimbursed is 100mg daily.

***MIDAZOLAM** (*1mg/mL, 5mg/mL Injection and generic brands*)

- For adjunctive therapy of pain management in palliative care patients outside the hospital setting. **[Criteria Code 01]**

***MIDOSTAURIN** (*Rydapt 25mg Capsule*)

- For the treatment of adult patients with newly diagnosed FMS-like tyrosine kinase 3 (FLT3)-mutated acute myeloid leukemia when used in combination with standard cytarabine and daunorubicin (7+3) induction and cytarabine consolidation chemotherapy. Patients should be deemed fit to receive standard induction and consolidation chemotherapy.

Clinical Notes:

- Midostaurin is not funded as maintenance therapy.
- Midostaurin may be used in combination with other 7+3 induction regimens (i.e. cytarabine and idarubicin)

Claim Note:

- Claims for Rydapt 25mg capsule that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00904390

MIGALASTAT (*Galafold 123mg Capsule*)

- Adults with confirmed diagnosis of Fabry Disease (alpha-galactosidase [alpha-Gal A]) and who have an alpha-Gal A mutation, determined to be amenable by an in vitro assay; and
- For use in patients with an amenable mutation and who are otherwise eligible for enzyme replacement therapy (ERT) for the treatment of Fabry Disease as determined through the Canadian Fabry Disease Initiative (CFDI).
- Not for use in pediatrics (i.e. patients < 18 years of age).

Clinical Note:

- Galafold will not be used concomitantly with any ERT.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claims Note:

- Claims for Galafold 123mg capsule that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904406
 - 00904407

MIRABEGRON (*Myrbetriq 25mg ER and 50mg ER Tablet*)

- See [OAB Medications](#)

MODAFINIL (*Alertec 100mg Tablet and generic brands*)

- For the treatment of narcolepsy confirmed by sleep study

***MOXIFLOXACIN** (*Avelox 400mg Tablet and generic brands*)

- See [Fluoroquinolones, Respiratory](#)

MOXIFLOXACIN (*Vigamox 0.5% Ophthalmic Solution and generic brands*)

- See [Fluoroquinolones, Ophthalmic](#)

NARATRIPTAN (*Amerge 1mg, 2.5mg Tablet and generic brands*)

- See [Selective 5HT₁ - Receptor Agonists](#)

NATALIZUMAB (*Tysabri 300mg/15mL Vial*)**Initial Request:**

For the treatment of Relapsing-Remitting Multiple Sclerosis (RRMS) who meet all the following criteria:

- The patient's physician is a neurologist experienced in the management of relapsing-remitting multiple sclerosis (RRMS); AND
 - The patient;
 - has a current EDSS less than or equal to 5.0; AND
 - has failed to respond to a full and adequate course¹ (at least six months) of at least ONE disease modifying therapy OR has contraindications/intolerance to at least TWO disease modifying therapies; AND
 - has had ONE of the following types of relapses in the past year:
 - the occurrence of one relapse with partial recovery during the past year AND has at least ONE gadolinium-enhancing lesion on brain MRI, OR significant increase in T2 lesion load compared to a previous MRI; OR
 - the occurrence of two or more relapses with partial recovery during the past year; OR
 - the occurrence of two or more relapses with complete recovery during the past year AND has at least ONE gadolinium-enhancing lesion on brain MRI, OR significant increase in T2 lesion load compared to a previous MRI.
- approval period: 1 year.

Requirements for Initial Requests:

- the patient's physician provides documentation setting out the details of the patient's most recent neurological examination within ninety (90) days of the submitted request. This must include a description of any recent attacks, the dates, and the neurological findings.
- MRI reports do NOT need to be submitted with the initial request.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Renewal:

- Date and details of the most recent neurological examination and EDSS scores must be provided (exam must have occurred within the last 90 days); AND
- Patients must be stable or have experienced no more than 1 disabling attack/relapse in the past year; AND
- Recent Expanded Disability Status Scale (EDSS) score less than or equal to 5.0.

1. Failure to respond to a full and adequate course is defined as a trial of at least one approved first line therapy for a minimum of 6 months AND experienced at least one disabling relapse (attack) while on this

***NETUPITANT/PALONOSETRON (*Akynzeo 300mg/0.5mg Capsule*)**

- In combination with dexamethasone for the prevention of acute and delayed nausea and vomiting in patients receiving:
 - highly emetogenic chemotherapy, **[Criteria Code 01]** OR
 - moderately emetogenic chemotherapy who have had inadequate symptom control using a 5-HT3 antagonist and dexamethasone in a previous cycle. **[Criteria Code 02]**

Clinical Notes:

- Highly emetogenic chemotherapy (HEC) may include, but is not limited to: cisplatin regimens, anthracycline and cyclophosphamide combination regimens, and regimens containing carmustine, mechlorethamine, streptozocin, dacarbazine and cyclophosphamide $\geq 1500\text{mg/m}^2$.
- Patients who receive carboplatin-based regimens with $\text{AUC} \geq 4$ are also eligible to receive netupitant/palonosetron in combination with dexamethasone for primary prevention of acute and delayed nausea and vomiting.

NILOTINIB (*Tasigna 150mg & 200mg Capsule*)*First Line:**

- For the first-line treatment of adult patients with Philadelphia chromosome positive chronic myeloid leukemia (Ph+ CML) in chronic phase.

Second Line:

- For the treatment of chronic phase and accelerated phase Philadelphia chromosome positive (Ph+) chronic myeloid leukemia (CML) in adult patients who:
 - Are resistant to imatinib;
 - Have progressed to accelerated phase while on imatinib;
 - Are intolerant to previous oral tyrosine kinase inhibitors (TKIs) (i.e. imatinib or dasatinib or both). Sequential use of nilotinib and dasatinib is not permitted except in cases of intolerance (i.e. grade 3 or 4 toxicity).

NINTEDANIB (*Ofev 100mg and 150mg Capsule*)**IDIOPATHIC PULMONARY FIBROSIS (IPF)**

- **Initial approval criteria:**
 - Adult patients who have a diagnosis of mild to moderate idiopathic pulmonary fibrosis (IPF)¹ confirmed by a respirologist and a high-resolution CT scan within the previous 24 months;
 - All other causes of restrictive lung disease (e.g. collagen vascular disorder or hypersensitivity pneumonitis) should be excluded;
 - Patient is under the care of a physician with experience in IPF;
 - Initial approval period: 7 months (allow 4 weeks for repeat pulmonary function tests).

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- **Initial renewal criteria:**
 - Patients must NOT demonstrate progression of disease defined as an absolute decline in percent predicted FVC of $\geq 10\%$ from initiation of therapy until renewal (initial 6 month treatment period). If a patient has experienced progression as defined above, then the results should be validated with a confirmatory pulmonary function test conducted 4 weeks later.
 - Approval period: 6 months.
- **Second and Subsequent renewal criteria (at 12 months after initiation and thereafter):**
 - Patients must NOT demonstrate progression of disease defined as an absolute decline in percent predicted FVC of $\geq 10\%$ within any 12 month period. If a patient has experienced progression as defined above, then the results should be validated with a confirmatory pulmonary function test conducted 4 weeks later.
 - Approval period: 12 months.
- **Exclusion Criteria:**
 - Combination use of Ofev (nintedanib) and Esbriet (pirfenidone) will not be funded.

1. Mild-moderate IPF is defined as: a forced vital capacity (FVC) $\geq 50\%$ of predicted.

Note:

- Patients who have experienced intolerance or failure to Ofev (nintedanib) or Esbriet (pirfenidone) will be considered for the alternate agent provided that the patient continues to meet the above coverage criteria.

CHRONIC FIBROSING INTERSTITIAL LUNG DISEASE

Initiation criteria:

- For the treatment of chronic fibrosing interstitial lung disease with a progressive phenotype confirmed by a specialist in interstitial lung diseases, if the following criteria are met:
 - the patient has a forced vital capacity greater than or equal to 45% of predicted.

Renewal criteria:

- The patient must not experience a more severe progression of disease, defined as an absolute decline in percent predicted forced vital capacity of 10% or greater over the preceding year of treatment with nintedanib.

Clinical Note:

- The patient's clinical status should be evaluated every 12 months.

Claim Notes:

- The patient is under the care of a physician with experience in interstitial lung diseases.
- Concurrent treatment of nintedanib with pirfenidone should not be reimbursed.
- Approval Period: 12 months

***NIRAPARIB (Zejula 100mg Tablet)**

NEWLY DIAGNOSED ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER

- As monotherapy maintenance treatment of patients with newly-diagnosed ovarian, fallopian tube, or primary peritoneal cancer who are in response (complete or partial) to at least 4 cycles of first-line platinum-based chemotherapy. Eligible patients should have high-grade serous or endometrioid tumours classified as stage III or IV according to the International Federation of Gynecology and Obstetrics (FIGO) criteria.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

- Patients should have a good performance status.
- Maintenance therapy with niraparib should begin within 12 weeks of completion of platinum- based chemotherapy and may continue for up to 3 years, or until disease progression or unacceptable toxicity, whichever occurs first.
- Patients who have stable brain metastases are eligible for treatment with niraparib.
- Patients who are unable to tolerate platinum-based chemotherapy (due to allergic reaction) and otherwise meet criteria, will be assessed on a case by case basis to determine eligibility for treatment with niraparib.
- Niraparib in combination with bevacizumab is not funded.

RELAPSED, PLATINUM SENSITIVE ADVANCED EPITHELIAL OVARIAN, FALLOPIAN TUBE OR PRIMARY PERITONEAL CANCER

- As monotherapy maintenance treatment for patients with relapsed, platinum-sensitive high grade serous epithelial ovarian, fallopian tube, or primary peritoneal cancer who have completed at least two previous lines of platinum-based chemotherapy, and have achieved a complete or partial response to the most recent platinum-based chemotherapy regimen.

Clinical Notes:

- Platinum-sensitive disease is defined as disease progression occurring at least six months after completion of platinum-based chemotherapy.
- Patients should have a good performance status.
- Patients must have received at least 4 cycles of the most recent platinum-based chemotherapy before starting treatment with niraparib.
- Maintenance therapy with niraparib should begin within 12 weeks of the last chemotherapy treatment and may continue until disease progression or unacceptable toxicity, whichever occurs first.
- Patients who have stable brain metastases are eligible for treatment with niraparib.
- Patients who are unable to tolerate platinum-based chemotherapy (due to allergic reaction) and otherwise meet criteria, will be assessed on a case by case basis to determine eligibility for treatment with niraparib.

Claim Note:

- Claims for Zejula 100mg tablet that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00904985

NITISINONE (Cycle-Nitisinone 2mg, 5mg and 10mg Tablet and Orfadin 2mg, 5mg, 10mg, 20mg Capsule)

- For the treatment of adult and pediatric patients with hereditary tyrosinemia type 1 (HT-1) in combination with dietary restriction of tyrosine and phenylalanine.

Clinical Note:

- For use in patients with an established diagnosis of HT-1.

Claim Notes:

- Must be prescribed by a physician experienced in the diagnosis and management of HT-1.
- Claims for nitisinone 10mg tablet/capsule and 20mg capsule that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - Nitisinone 10mg Tab

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- 00904442
- 00904443
- 00904444
- Orfadin 10mg Cap
 - 00904434
 - 00904435
 - 00904436
- Orfadin 20mg Cap
 - 00904437
 - 00904438
 - 00904439

***NORFLOXACIN (400mg Tablet and generic brands)**

- See [Fluoroquinolones, Oral](#)

NUSINERSEN (Spinraza 12mg/5mL Vial)

- For patients diagnosed with 5q Spinal Muscular Atrophy (SMA) under the care of a specialist with experience in the diagnosis and management of SMA, if the following clinical criteria are met:
 - Genetic documentation of 5q SMA homozygous gene deletion, homozygous mutation, or compound heterozygote, AND
 - Patients who:
 - are pre-symptomatic with two or three copies of SMN2, OR
 - have had disease duration of less than six months, two copies of SMN2, and symptom onset after the first week after birth and on or before seven months of age, OR
 - are under the age of 18 with symptom onset after six months of age,
 AND
 - Patient is not currently requiring permanent invasive ventilation*, AND
 - A baseline assessment using an age-appropriate scale (the Hammersmith Infant Neurological Examination [HINE] Section 2, Children’s Hospital of Philadelphia Infant Test of Neuromuscular Disorders [CHOP INTEND], or Hammersmith Functional Motor Scale-Expanded [HFMSE]) must be completed prior to initiation of nusinersen treatment.
- For continued coverage, the patient must meet the following criteria:
 - There is demonstrated achievement or maintenance of motor milestone function (as assessed using age-appropriate scales: the [HINE] Section 2), CHOP INTEND, or HFMSE) since treatment initiation in patients who were pre-symptomatic at the time of treatment initiation; OR
 - There is demonstrated maintenance of motor milestone function (as assessed using age-appropriate scales: the HINE Section 2, CHOP INTEND, or HFMSE) since treatment initiation in patients who were symptomatic at the time of treatment initiation;
 AND
 - Patient does not require permanent invasive ventilation*.
- Treatment should be discontinued if, prior to the fifth dose or every subsequent dose of nusinersen, the above renewal criteria are not met.

* Permanent invasive ventilation is defined as the use of tracheostomy and a ventilator due to progression of SMA that is not due to an

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

identifiable and reversible cause.

Claim Note:

- Coverage for nusinersen will not be provided in combination with other SMA drug therapies or post administration of onasemnogene abeparvovec.
- Claims for Spinraza 12mg/5mg vials that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904366
 - 00904367
 - 00904368
 - 00904369
 - 00904370
 - 00904371
 - 00904372
 - 00904373
 - 00904374
 - 00904375
 - 00904376
 - 00904377
 - 00904474
 - 00904475
 - 00904476
 - 00904478

OAB MEDICATIONS (*Darifenacin, Fesoterodine Fumarate, Mirabegron, and Trospium*)

- For the treatment of overactive bladder (OAB) with symptoms of urgency, urgency incontinence, and urinary frequency in patients who have an intolerance or insufficient response to an adequate trial of immediate-release oxybutynin, solifenacin or tolterodine.

OBETICHOLIC ACID (*Ocaliva 5mg and 10mg Tablet*)

Initiation Criteria:

- For the treatment of primary biliary cholangitis (PBC) in combination with ursodeoxycholic acid (UDCA) in adults with an inadequate response to UDCA, or as monotherapy in adults unable to tolerate UDCA, where the following criteria are met:
 - A confirmed diagnosis of PBC, defined as:
 - Positive antimitochondrial antibodies (AMA); **or**
 - Liver biopsy results consistent with PBC.
 - The patient is under the care of a gastroenterologist or hepatologist or other prescriber with a specialty in gastroenterology or hepatology.
- AND**
- The patient has received UDCA for a minimum of 12 months and has experienced an inadequate response to UDCA and can benefit from the addition of obeticholic acid. An inadequate response is defined as:

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- alkaline phosphatase (ALP) ≥ 1.67 x upper limit of normal (ULN) **and/or**
- bilirubin $> \text{ULN}$ and $< 2 \times \text{ULN}$ **and/or**
- evidence of compensated cirrhosis

OR

- The patient has experienced documented and unmanageable intolerance to UDCA and can benefit from switching therapy to obeticholic acid.

Renewal Criteria:

- The patient continues to benefit from treatment with obeticholic acid as evidenced by:
 - A reduction in the ALP level to less than $1.67 \times \text{ULN}$; **or**
 - A 15% reduction in the ALP level compared with values before beginning treatment with obeticholic acid.

Claim Note:

- Duration of approval: 12 months

OCRELIZUMAB (*Ocrevus 300mg/10mL Vial*)

PRIMARY PROGRESSIVE MULTIPLE SCLEROSIS

- For the treatment of adult patients with early primary progressive multiple sclerosis (PPMS) who meet all of the following criteria:
 - Confirmed diagnosis based on McDonald criteria
 - Recent Expanded Disability Status Scale (EDSS) score between 3.0 and 6.5
 - Recent Functional Systems Scale (FSS) score of at least 2 for the pyramidal functions component due to lower extremity findings
 - Disease duration of 10 years for those with an EDSS of less than or equal to 5 or disease duration less than 15 years for those with an EDSS greater than 5
 - Diagnostic imaging features characteristic of inflammatory activity
 - Must be prescribed by a neurologist with experience in the diagnosis and management of multiple sclerosis.

Clinical Note:

- Treatment should be discontinued for patients with an EDSS score of greater than or equal to 7.

RELAPSING REMITTING MULTIPLE SCLEROSIS

- For the treatment of adult patients with relapsing remitting multiple sclerosis (RRMS) who meet all of the following criteria:
 - Confirmed diagnosis based on McDonald criteria
 - Experienced one or more disabling relapses or new MRI activity in the last two years
 - Are fully ambulatory without aids (i.e., must provide a recent Expanded Disability Status Scale (EDSS) score of less than or equal to 5.5)
 - Must be prescribed by a neurologist with experience in the diagnosis and management of multiple sclerosis.

Clinical Note:

- Treatment should be discontinued for patients with an EDSS score of greater than or equal to 6.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Notes:

- Combined use with other disease modifying therapies to treat RRMS will not be reimbursed.
- Claims for Ocrevus 300mg/10mL Vial that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00904527

OFATUMUMAB (*Kesimpta 20mg/0.4mL Prefilled Pen*)**RELAPSING REMITTING MULTIPLE SCLEROSIS (RRMS)**

- For the treatment of adult patients with relapsing remitting multiple sclerosis (RRMS) who meet all of the following criteria:
 - An Expanded Disability Status Scale (EDSS) score of less than 6.0
 - Evidence of active disease defined as at least one of the following:
 - One relapse during the previous year
 - Two relapses during the previous 2 years
 - A positive gadolinium (Gd)-enhancing MRI scan during the year before starting treatment with ofatumumab.

Renewal Criteria:

- EDSS score less than 6.0. Date and details of the most recent neurological examination and EDSS score must be provided (exam must have occurred within the last 90 days);
AND
- Patients must be stable or have experienced no more than 1 disabling attack/relapse in the past year.

Claims Notes:

- Approval: 1 year.
- Combined use with other disease modifying therapies to treat multiple sclerosis will not be reimbursed.
- Must be prescribed by a neurologist with experience in the diagnosis and management of multiple sclerosis.

OFLOXACIN, OPHTHALMIC (*Ocuflox 0.3% Ophthalmic Solution and generic brands*)

- See [Fluoroquinolones, Ophthalmic](#)

***OLAPARIB** (*Lynparza 50mg Capsule and 100mg and 150mg Tablet*)**HIGH-RISK EARLY BREAST CANCER**

For the treatment of adult patients with germline BRCA-mutated (gBRCAm) HER2-negative high-risk early breast cancer who meet one of the following criteria:

- For patients who underwent upfront surgery followed by adjuvant chemotherapy:
 - If TNBC: must have node-positive disease or pT ≥ 2cm, or
 - If HR-positive, HER2-negative: must have ≥ 4 involved pathologically confirmed lymph nodesOR
- For patients who received neoadjuvant chemotherapy followed by surgery:
 - If TNBC: must have residual invasive disease, or
 - If HR-positive, HER2-negative: must have residual invasive disease and deemed high-risk using a risk assessment tool.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

- Patients should have a good performance status.
- Treatment should continue until disease recurrence, unacceptable toxicity, or to a maximum of one year, whichever occurs first.
- Must have confirmed gBRCAm prior to starting therapy.
- Patients must have completed chemotherapy containing anthracyclines and/or taxanes. Patients who stop chemotherapy early for toxicity are eligible.

NEWLY-DIAGNOSED, ADVANCED, BRCA-MUTATED

- As monotherapy maintenance treatment of patients with newly-diagnosed, advanced, BRCA-mutated (germline or somatic), high-grade epithelial ovarian, fallopian tube, or primary peritoneal cancer who are in response (complete or partial) to at least 4 cycles of first-line platinum-based chemotherapy.

Clinical Notes:

- Patients should have a good performance status.
- Maintenance therapy with olaparib should begin within 12 weeks of completion of platinum-based chemotherapy.
- Patients who are unable to tolerate platinum-based chemotherapy (due to allergic reaction) and otherwise meet criteria, will be assessed on a case by case basis to determine eligibility for treatment with olaparib.
- Treatment should continue until unacceptable toxicity, disease progression, or to a maximum of 2 years of therapy if no evidence of disease, whichever comes first.¹
- Imaging is required for patients who are delayed in starting olaparib therapy, i.e. greater than 12 weeks after completion of platinum-based chemotherapy, or who have had a break in therapy for more than 14 days, to rule out progression prior to starting or re-starting olaparib.
- Olaparib in combination with bevacizumab is not funded. Patients already on bevacizumab maintenance at the time of olaparib funding may be switched to olaparib, as long as there is no evidence of progression on imaging and is within 12 weeks of completion of chemotherapy.

¹. Patients with a partial response or stable disease at 2 years may continue to receive olaparib at the discretion of the treating physician.

RELAPSED, BRCA-MUTATED

- As monotherapy maintenance treatment for patients with platinum-sensitive, relapsed, BRCA-mutated (germline or somatic), high grade serous epithelial ovarian, fallopian tube, or primary peritoneal cancer who have completed at least two previous lines of platinum-based chemotherapy and are in radiologic response (complete or partial) to their most recent platinum-based chemotherapy regimen as per the SOLO-2 trial.
- Patients must have received at least four cycles of their most recent platinum-based chemotherapy before starting treatment with olaparib.

Clinical Notes:

- Treatment should continue until unacceptable toxicity or disease progression.
- Maintenance therapy with olaparib should begin within eight weeks of the last dose of platinum-based chemotherapy.
- Platinum-sensitive disease is defined as disease progression occurring at least six months after completion of platinum-based chemotherapy.
- Patients should have a good performance status.
- Patients who are unable to tolerate platinum-based chemotherapy (due to allergic reaction) and otherwise meet criteria, will be assessed on a case by case basis to determine eligibility for treatment with olaparib.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

METASTATIC CASTRATE-RESISTANT PROSTATE CANCER

- For the treatment of patients with metastatic castration-resistant prostate cancer (mCRPC) with deleterious or suspected deleterious germline and/or somatic mutations in the homologous recombination repair (HRR) genes BRCA1, BRCA2 or ATM and who have progressed on prior treatment with androgen-receptor-axis-targeted (ARAT) therapy.

Clinical Note:

- Patients should have a good performance status and treatment should be continued until disease progression or unacceptable toxicity.

OMALIZUMAB (*Xolair 150mg Powder for Injection and Prefilled Syringe*)

- For the treatment of adults and adolescents (12 years of age or older) with moderate to severe chronic idiopathic urticaria (CIU) who remain symptomatic (presence of hives and/or associated itching) despite optimum management with available oral therapies.

Criteria Notes:

- Prescribed by a specialist (allergist, immunologist, dermatologist, etc.) or other authorized prescriber with knowledge of CIU treatment.
- Initial approval period of 24 weeks at a maximum dose of 300mg every 4 weeks.
- Treatment cessation could be considered for patients who experience complete symptom control for at least 12 consecutive weeks at the end of a 24 week treatment period.
- Continued coverage will be authorized if the patient has achieved:
 - complete symptom control for less than 12 consecutive weeks; or
 - partial response to treatment, defined as at least a ≥ 9.5 point reduction in baseline urticaria activity score over 7 days (UAS7)

ONABOTULINUMTOXIN-A (*Botox 50iu/vial and 100iu/vial Injection*)

- For the treatment of the following Health Canada approved indications:
 - focal spasticity following stroke in adults;
 - equinus foot deformity in cerebral palsy patients 2 years of age and older;
 - cervical dystonia;
 - blepharospasm, hemifacial spasm (VII nerve disorder) or strabismus in patients 12 years of age and older.

ONABOTULINUMTOXIN-A (*Botox 50iu/vial and 100iu/vial Injection*)

- For the treatment of overactive bladder (OAB) with symptoms of urgency, urgency incontinence, and urinary frequency, in adult patients who have an intolerance or insufficient response to an adequate trial of at least two other pharmacologic treatments (e.g. anticholinergics, mirabegron).

Renewal criteria:

- Requests for renewal should provide objective evidence of a treatment response, defined as a reduction of at least 50% in the frequency of urinary incontinence episodes.

Claim Notes:

- Must be prescribed and administered by a urologist.
- Initial approval period: 12 weeks (one dose).

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Renewal approval period: Maximum of 3 doses per year in responders, at a frequency of no more than once every twelve weeks.

ONABOTULINUMTOXIN-A (*Botox 200iu/vial Injection*)

- For the treatment of urinary incontinence due to neurogenic detrusor overactivity resulting from neurogenic bladder associated with multiple sclerosis (MS) or subcervical spinal cord injury (SCI) in patients who have failed to respond to behavioural modification and anticholinergics and/or are intolerant to anticholinergics.
- Subsequent treatments are provided at intervals no less than every 36 weeks.

***ONDANSETRON** (*Zofran 4mg, 8mg Tablet, 4mg/5mL Oral Liquid, ODT Tablet and generic brands*)

- See [Serotonin \(5-HT₃\) Antagonists](#)

Note:

- Only requests for the oral dosage forms are eligible for consideration. Although the dose may vary, usually a single oral 8mg dose pre-chemotherapy is sufficient to control symptoms. As well, some patients may require additional therapy up to 48 hours after the last dose of chemotherapy or last radiation treatment. Benefit beyond 48 hours has not been established and is therefore, not insured.

***OSELTAMIVIR** (*Tamiflu 30mg, 45mg, 75mg Capsule, 6mg/mL Oral Suspension and generic brands*)

Treatment: [Criteria Code 40]

- For patients who test negative for COVID-19 and meet one of the following:
 1. have suspected¹ or test confirmed severe, complicated, or progressive² influenza OR
 2. are hospitalized² with suspected¹ or test confirmed influenza OR
 3. have suspected¹ or test confirmed influenza and are at higher risk of complications, which include the following age groups, chronic medical conditions, and persons:
 - Asthma and other chronic pulmonary disease, including asthma, bronchopulmonary dysplasia, cystic fibrosis, chronic bronchitis, and emphysema
 - Cardiovascular disease (excluding isolated hypertension; including congenital and acquired heart disease, such as congestive heart failure and symptomatic coronary artery disease)
 - Renal disease
 - Chronic liver disease
 - Diabetes mellitus and other metabolic diseases
 - Anemia and hemoglobinopathies, such as sickle cell disease
 - Cancer, immunosuppression, or immunodeficiency due to disease (e.g.: HIV infection, especially if CD4 is <200) or management of underlying condition (solid organ transplant or hematopoietic stem cell transplant recipients)
 - Neurological disease and neurodevelopmental disorders that compromise handling of respiratory secretions (cognitive dysfunction; spinal cord injury; neuromuscular, neurovascular, neurodegenerative, and seizure disorders; cerebral palsy; metabolic disorders)
 - Children aged younger than 5 years²
 - Individuals aged 65 years or older
 - People of any age who are residents of nursing homes or other chronic care facilities
 - Pregnancy and up to 4 weeks postpartum regardless of how the pregnancy ended³
 - Obesity with a BMI ≥40 or a BMI >3 z-scores above the mean for age and gender
 - Children and adolescents aged younger than 18 years undergoing treatment for long periods with

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

acetylsalicylic acid because of the potential increase in Reye's syndrome associated with influenza

- Indigenous peoples
- For the treatment of long-term care and eligible⁴ residential care residents with clinically suspected or lab confirmed influenza A or B upon advice of the Medical Officer of Health. A clinically suspected case is one in which the patient meets the criteria of influenza-like illness and there is confirmation of influenza A or B circulating within the facility or surrounding community.

Prophylaxis: [Criteria Code 41]

- For the prophylaxis of influenza A or B in long-term care and eligible⁴ residential care residents where the facility has an outbreak upon advice of the Medical Officer of Health.
- A protocol has been developed by Public Health for the treatment of residents in long-term care facilities and eligible⁴ residential care residents. The facility must contact the Medical Officer of Health or local Public Health Office who will notify the Pharmacare office (or dispensing pharmacy after office hours) if coverage is required.

Notes:

1. For suspected cases, discontinue oseltamivir if the lab test is negative
2. Among healthy children aged younger than 5 years, the risk of hospitalization is further increased among those aged younger than 2 years
3. The risk of influenza-related hospitalization increases with length of gestation (i.e., it is higher in the third trimester than in the second)
4. Eligible residents are people of any age who are residents of nursing homes or other chronic care facilities.

***OSIMERTINIB (*Tagrisso 40mg and 80mg Tablet*)**

STAGE IB-III A NON-SMALL CELL LUNG CANCER (NSCLC)

- For adjuvant therapy after tumour resection in patients with Stage IB-III A (AJCC 7th edition or equivalent) non-small cell lung cancer (NSCLC) whose tumors have epidermal growth factor receptor (EGFR) exon 19 deletions [exon 19 del] or exon 21 [L858R] substitution mutations.

Clinical Notes:

- Patients should have a good performance status.
- Treatment with osimertinib should continue for a total duration of 3 years, or until disease recurrence or unacceptable toxicity.
- Osimertinib treatment should be initiated within 10 weeks of complete surgical resection if adjuvant chemotherapy was not administered, or within 26 weeks if adjuvant chemotherapy was administered.
- Retreatment with osimertinib in the metastatic setting will be considered if disease recurrence is at least 6 months following completion of adjuvant therapy.

LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER (NSCLC)

- For the first-line treatment of patients with locally advanced (not amenable to curative-intent therapy) or metastatic non-small cell lung cancer (NSCLC) whose tumors have the following epidermal growth factor receptor (EGFR) mutations: exon 19 deletions [exon 19 del] or exon 21 [L858R] mutations. Eligible patients should be previously untreated in the locally advanced or metastatic setting and have a good performance status.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

- Treatment may continue until clinically meaningful disease progression or unacceptable toxicity.
- Retreatment with osimertinib in the metastatic setting will be considered if disease recurrence is at least 6 months following completion of adjuvant therapy.

LOCALLY ADVANCED OR METASTATIC T790M MUTATION-POSITIVE NON-SMALL CELL LUNG CANCER (NSCLC)

- For the treatment of patients with locally advanced or metastatic epidermal growth factor receptor (EGFR) T790M mutation-positive non-small cell lung cancer (NSCLC) who have progressed on EGFR tyrosine kinase inhibitor (TKI) therapy, or as initial therapy in patients with a de novo EGFR T790M mutation.

Clinical Notes:

- Patients currently receiving alternate first-line EGFR TKI's (e.g. erlotinib, gefitinib, afatinib) whose tumors have the noted EGFR mutations (exon 19 del or L858R) may be switched to osimertinib provided they meet all other funding criteria and have not experienced disease progression.
- Patients who have initiated treatment with chemotherapy prior to receiving results of the EGFR mutation status may be switched to osimertinib if otherwise eligible.
- Osimertinib may be continued until there is evidence of disease progression or the development of unacceptable toxicity.
- Retreatment with osimertinib in the metastatic setting will be considered if disease recurrence is at least 6 months following completion of adjuvant therapy.

OXCARBAZEPINE (*Trileptal 60mg/mL Oral Liquid and 150mg, 300mg, 600mg Tablet and generic brands*)

- For the treatment of epileptic seizures in patients who have had an inadequate response to or are intolerant of at least three other formulary agents (prior or current use) including carbamazepine.

OZANIMOD (*Zeposia 0.23mg & 0.46mg Initiation Pack and 0.92mg Cap*)

- For the treatment of adult patients with moderately to severely active ulcerative colitis (UC) who have a partial Mayo score > 4, and have a rectal bleeding subscore ≥ 2, and are:
 - refractory or intolerant to conventional therapy (i.e. 5-ASA for a minimum of 4 weeks, and prednisone ≥ 40mg daily for two weeks or IV equivalent for one week); OR
 - corticosteroid dependent (i.e. cannot be tapered from corticosteroids without disease recurrence; or have relapsed within three months of stopping corticosteroids; or require two or more courses of corticosteroids within one year.)
- Renewal requests must include information demonstrating the beneficial effects of the treatment, specifically:
 - a decrease in the partial Mayo score ≥ 2 from baseline, AND
 - a decrease in the rectal bleeding subscore ≥ 1.

Clinical Notes:

- 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. The nature of intolerance(s) must be clearly documented.
- Patients with severe disease do not require a trial of 5-ASA.

Claim Notes:

- Must be prescribed by a gastroenterologist or physician with a specialty in gastroenterology.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Concurrent use of biologics or Janus kinase inhibitors not approved.
- Initial Approval: 16 weeks.
- Maximum dose of 0.92mg daily with no dose escalation permitted.
- Renewal Approval: 1 year.

***PALBOCICLIB** (*Ibrance 75mg, 100mg and 125mg Capsule and 75mg, 100mg and 125mg Tablet*)

ER POSITIVE, HER2-NEGATIVE ADVANCED BREAST CANCER IN COMBINATION WITH AN AROMATASE INHIBITOR (AI)

- In combination with an aromatase inhibitor (AI) (i.e. letrozole, anastrozole or exemestane) for the treatment of post-menopausal women with estrogen receptor (ER) positive, human epidermal growth factor receptor 2 (HER 2) negative advanced breast cancer who have not received any prior endocrine-based treatment for metastatic disease. Patients may have received up to one prior line of chemotherapy for advanced disease.

Clinical Notes:

- Treatment should continue until unacceptable toxicity or disease progression.
- Patients should have a good performance status and not be resistant to prior (neo) adjuvant aromatase inhibitor therapy (i.e. have the potential to benefit from first-line endocrine based therapy), without active or uncontrolled metastases to the central nervous system.
- Patients will be eligible for either palbociclib plus an aromatase inhibitor in the first line setting or everolimus plus exemestane as a subsequent line of therapy, but not both therapies. Patients eligible include:
 - Pre and peri-menopausal patients (should be treated with a luteinizing hormone- releasing hormone (LHRH) agonist)
 - Males
 - Patients with bone-only metastases
 - Patients who are HER2 equivocal by FISH testing (these patients are HER2 negative)
 - Patients currently receiving first line aromatase inhibitor monotherapy for ER positive, HER2-negative metastatic breast cancer may have palbociclib added provided the above criteria is met.
- Patients with prior adjuvant treatment with abemaciclib plus endocrine therapy are eligible if disease progression occurred 6 months or greater after completion of adjuvant abemaciclib.

HR POSTIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION WITH FULVESTRANT

- In combination with fulvestrant for the treatment of patients with hormone receptor (HR) positive, HER 2 negative advanced or metastatic breast cancer, as initial endocrine-based therapy or following disease progression on endocrine therapy. Patients may have also received up to one prior line of chemotherapy for advanced disease. Patients should have a good performance status, without active or uncontrolled metastases to the central nervous system and can be of any menopausal status (Perimenopausal and premenopausal women must be treated with an LHRH agonist).

Clinical Notes:

- Treatment should continue until unacceptable toxicity or disease progression.
- Patients who progress ≤ 12 months from (neo) adjuvant therapy are eligible for treatment with palbociclib plus fulvestrant.
- Patients who experience disease progression on prior CDK 4/6 inhibitor therapy, fulvestrant or everolimus are not eligible for treatment with palbociclib with fulvestrant.
- Patients currently receiving fulvestrant monotherapy, and who have not progressed may have palbociclib added, provided they are CDK 4/6 inhibitor naïve and otherwise meet funding criteria.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Patients who previously received everolimus plus exemestane will be eligible for funding of palbociclib plus fulvestrant on progression, provided that treatment was started prior to funding of CDK 4/6 + fulvestrant, patient must be CDK 4/6 naïve and otherwise meet funding criteria.
- Patients with prior adjuvant treatment with abemaciclib plus endocrine therapy are eligible if disease progression occurred 6 months or greater after completion of adjuvant abemaciclib.

PALIPERIDONE (*Invega Sustenna 50mg/0.5mL, 75mg/0.75mL, 100mg/mL and 150mg/1.5mL Injection Kit*)

- For the maintenance treatment of schizophrenia and related psychotic disorders (not dementia related) in patients who are not adherent to an oral antipsychotic; OR
- Who are currently receiving a long-acting injectable antipsychotic and require an alternative long acting injectable antipsychotic.

PALIPERIDONE (*Invega Trinza 175mg/0.875mL, 263mg/1.315mL, 350mg/1.75mL and 525 mg/2.625mL Injection*)

- For the maintenance treatment of schizophrenia and related psychotic disorders (not dementia related) in patients who have been stabilized on therapy with injectable paliperidone for at least four months.

***PANTOPRAZOLE MAGNESIUM** (*Tecta 40mg Tablet and generic brands*)

- See [Proton Pump Inhibitors](#)

PATISIRAN (*Onpattro 2mg/mL IV Solution*)

- For the treatment of polyneuropathy in adult patients with hereditary transthyretin-mediated amyloidosis (hATTR) who meet all of the following criteria:
 - Confirmed genetic diagnosis of hATTR.
 - Symptomatic with early-stage neuropathy¹.
 - Does not have New York Heart Association class III or IV heart failure.
 - Has not previously undergone a liver transplant.

Discontinuation Criteria:

- The patient is permanently bedridden and dependent on assistance for basic activities of daily living.
- OR**
- The patient is receiving end-of-life care.

Clinical Note:

1. Symptomatic early-stage neuropathy is defined as polyneuropathy disability stage I to IIIB or familial amyloidotic polyneuropathy stage I or II.

Claim Notes:

- The patient must be under the care of a physician with experience in the diagnosis and management of hATTR.
- Combination therapy with other interfering ribonucleic acid drugs or transthyretin stabilizers used to treat hATTR will not be reimbursed.
- Initial Approval: 9 months.
- Renewal Approval: 12 months. Confirmation of continued response is required.
- Claims for Onpattro 2mg/mL IV Solution that exceed the maximum claim amount of \$9,999.99 must be divided

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

and submitted as separate transactions using the DIN first and then the following PINs:

- 00904586
- 00904587
- 00904588

***PAZOPANIB** (*Votrient 200mg Tablet and generic brands*)

- For patients with advanced or metastatic renal cell carcinoma as either first-line therapy, or second-line therapy after failure of first-line immunotherapy.

Clinical Notes:

- Patients must have a good performance status.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.
- Pazopanib may not be used after another tyrosine kinase inhibitor (i.e., sorafenib, or sunitinib) as sequential therapy.
- In the event of significant toxicity, a switch to another tyrosine kinase inhibitor (i.e., sorafenib or sunitinib) may be allowed.
- Both clear cell and non-clear cell histology are eligible for treatment.

PDP-AMLODIPINE (*1mg/mL Oral Solution*)

- For patients who require administration through a feeding tube. **[Criteria Code 37]**
- For patients 19 years of age and younger, who cannot use a tablet or capsule. **[Criteria Code 38]**

***PEGFILGRASTIM** (*Fulphila 6mg/0.6mL (10mg/mL) PF Solution for Injection, Lapelga 6mg Prefilled Syringe, & 10mg/mL Prefilled Autoinjector, Nyvepria 10mg/mL Prefilled Syringe and Ziextenzo 10mg/mL Injection*)

- For the prevention of febrile neutropenia in patients with non-myeloid malignancies receiving myelosuppressive chemotherapy with curative intent who:
 - are at high risk of febrile neutropenia due to chemotherapy regimen, co-morbidities or pre-existing severe neutropenia; or
 - have had an episode of febrile neutropenia, neutropenic sepsis or profound neutropenia in a previous cycle of chemotherapy; or
 - have had a dose reduction, or treatment delay greater than one week due to neutropenia.

Clinical Note:

- Patients with non-curative cancer receiving chemotherapy with palliative intent are not eligible for coverage of pegfilgrastim for prevention of febrile neutropenia.

PERAMPANEL (*Fycompa 2mg, 4mg, 6mg, 8mg, 10mg and 12mg Tablet*)

- For the adjunctive treatment of refractory partial-onset seizures or primary generalized tonic-clonic seizures in patients who are currently receiving two or more antiepileptic drugs, and who have had an inadequate response to at least three other antiepileptic drugs.

Claim Note:

- The patient must be under the care of a physician experienced in the treatment of epilepsy.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

*PILOCARPINE, ORAL (*Salagen 5mg Tablet and generic brands*)

- For oncology patients only.
- For the treatment of the symptoms of xerostomia due to salivary gland hypofunction caused by radiotherapy for cancer of the head and neck.

PIRFENIDONE (*Esbriet 267mg Capsule, 267mg Tablet, 801mg Tablet and generic brands*)

- **Initial approval criteria:**
 - Adult patients who have a diagnosis of mild to moderate idiopathic pulmonary fibrosis (IPF)¹ confirmed by a respirologist and a high-resolution CT scan within the previous 24 months;
 - All other causes of restrictive lung disease (e.g. collagen vascular disorder or hypersensitivity pneumonitis) should be excluded;
 - Patient is under the case of a physician with experience in IPF;
 - Initial approval period: 7 months (allow 4 weeks for repeat pulmonary function tests).
 - **Initial renewal criteria:**
 - Patients must NOT demonstrate progression of disease defined as an absolute decline in percent predicted FVC of $\geq 10\%$ from initiation of therapy until renewal (initial 6 month treatment period). If a patient has experienced progression as defined above, then the results should be validated with a confirmatory pulmonary function test conducted 4 weeks later;
 - Approval period: 6 months
 - **Second and Subsequent renewal criteria (at 12 months after initiation and thereafter):**
 - Patients must NOT demonstrate progression of disease defined as an absolute decline in percent predicted FVC of $\geq 10\%$ within any 12 month period. If a patient has experienced progression as defined above, then the results should be validated with a confirmatory pulmonary function test conducted 4 weeks later;
 - Approval period: 12 months
 - **Exclusion Criteria:**
 - Combination use of Esbriet (pirfenidone) and Ofev (nintedanib) will not be funded.
1. Mild-moderate IPF is defined as: a forced vital capacity (FVC) $\geq 50\%$ of predicted.

*PLERIXAFOR (*Mozobil 24mg/1.2mL (20mg/mL) Single Use Vial and generic brands*)

- For use in combination with filgrastim to mobilize hematopoietic stem cells for subsequent autologous transplantation in patients who meet one of the following criteria:
 - PBCD34+ count of less than 10 cells/uL after 4 days of filgrastim, or
 - Less than 50% of the target CD34+ yield is achieved on the first day of apheresis (after being mobilized with filgrastim alone or following chemotherapy), or
 - Failed a previous attempt for stem cell mobilization with filgrastim alone or following chemotherapy.

Claim Note:

- Reimbursement is limited to a maximum of 4 doses (0.24mg/kg given daily) for a single mobilization attempt and to prescriptions written by an oncologist or hematologist.

*POMALIDOMIDE (*Pomalyst 1mg, 2mg, 3mg, and 4mg Capsule and generic brands*)

- For patients with relapsed and/or refractory multiple myeloma who have previously failed at least two treatments, including both bortezomib and lenalidomide and demonstrated disease progression on the last treatment.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Pomalidomide may be an option in rare instances where bortezomib is not tolerated or contraindicated but in all cases, patients should have failed lenalidomide.

Note:

- Pomalidomide must be prescribed and dispensed only by physicians and pharmacists who are registered with and agree in writing to adhere to the guidelines of the Company's RevAid® Program. Details are available at <https://revaaid.ca/revaaid>.

***PONATINIB (*Iclusig 15mg Tablet*)**

- For the treatment of patients with chronic phase, accelerated phase or blast phase chronic myeloid leukemia (CML) or Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL) for whom other tyrosine kinase inhibitor (TKI) therapy is not appropriate, including CML or Ph+ ALL that is T315i mutation positive or where there is resistance or intolerance to prior TKI therapy. Funding should be for ECOG performance status 0-2. Treatment should continue until unacceptable toxicity or disease progression.

PRASUGREL (*10mg Tablet*)

- In combination with ASA for patients with:
 - Unstable angina (UA) or non-ST-segment elevation myocardial infarction (NSTEMI) managed with percutaneous coronary intervention (PCI); or
 - ST-segment elevation myocardial infarction (STEMI) managed with primary or delayed PCI; or
 - Failure on clopidogrel and ASA therapy as defined by definite stent thrombosis, or recurrent STEMI, NSTEMI or UA after revascularization with PCI.

Clinical Note:

- Definite stent thrombosis, according to the Academic Research Consortium, is a total occlusion originating in or within 5 mm of the stent or is a visible thrombus within the stent or is within 5 mm of the stent in the presence of an acute ischemic clinical syndrome within 48 hours.

Claim Note:

- Approval Period: 1 year.

PROPIVERINE HYDROCHLORIDE (*Mictoryl 5mg Tablet*)

- For the treatment of overactive bladder with symptoms of urgency incontinence and/or urinary frequency and urgency in pediatric patients under 18 years of age.

PROPRANOLOL (*Hemangirol 3.75mg/mL Solution*)

- For the treatment of patients with proliferating infantile hemangioma that is:
 - Life-or function-threatening OR
 - Ulcerated with pain or not responding to simple wound care measures OR
 - At risk of permanent scarring or disfigurement

***PROTON PUMP INHIBITORS (PPIs)**

PANTOPRAZOLE MAGNESIUM AND LANSOPRAZOLE

- Failure of a trial of all open benefit PPIs (omeprazole, pantoprazole sodium and rabeprazole).

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

*REGORAFENIB (*Stivarga 40mg Tablet*)

GASTROINTESTINAL STROMAL TUMORS (GIST)

- For patients with metastatic and/or unresectable gastrointestinal stromal tumors (GIST) who have had disease progression on, or intolerance to, imatinib and sunitinib; AND has ECOG \leq 1.

HEPATOCELLULAR CARCINOMA (HCC)

- For the treatment of patients with unresectable hepatocellular carcinoma (HCC) who have experienced disease progression on sorafenib or lenvatinib and meet all of the following criteria:
 - Child-Pugh class status of A.
 - ECOG performance status of 0 or 1.

Clinical Notes:

- Treatment should continue until disease progression or unacceptable toxicity.
- Patients with disease progression on sorafenib must have tolerated a minimum dose of 400 mg per day for at least 20 of the last 28 days of treatment.

*RIBOCICLIB (*Kisqali 200mg Tablet*)

ER POSITIVE, HER2-NEGATIVE ADVANCED BREAST CANCER IN COMBINATION WITH AN AROMATASE INHIBITOR (AI)

- In combination with an aromatase inhibitor (AI) (i.e. letrozole, anastrozole or exemestane) for the treatment of post-menopausal women with estrogen receptor (ER) positive, human epidermal growth factor receptor 2 (HER 2) negative advanced breast cancer who have not received any prior endocrine-based treatment for metastatic disease. Patients may have received up to one prior line of chemotherapy for advanced disease.

Clinical Notes:

- Treatment should continue until unacceptable toxicity or disease progression.
- Patients should have a good performance status and not be resistant to prior (neo) adjuvant aromatase inhibitor therapy (i.e. have the potential to benefit from first-line endocrine based therapy), without active or uncontrolled metastases to the central nervous system.
- Patients will be eligible for either ribociclib plus an aromatase inhibitor in the first line setting or everolimus plus exemestane as a subsequent line of therapy, but not both therapies. Patients eligible include:
 - Pre and peri-menopausal patients (should be treated with a luteinizing hormone- releasing hormone (LHRH) agonist)
 - Males
 - Patients with bone only metastases
 - Patients who are HER2 equivocal by FISH testing (these patients are HER2 negative)
 - Patients currently receiving first line aromatase inhibitor monotherapy for ER positive, HER2-negative metastatic breast cancer may have ribociclib added provided the above criteria is met.
 - Patients with prior adjuvant treatment with abemaciclib plus endocrine therapy are eligible if disease progression occurred 6 months or greater after completion of adjuvant abemaciclib.

HR POSITIVE, HER2-NEGATIVE ADVANCED OR METASTATIC BREAST CANCER IN COMBINATION WITH FULVESTRANT

- In combination with fulvestrant for the treatment of patients with hormone receptor (HR) positive, HER 2 negative advanced or metastatic breast cancer, as initial endocrine-based therapy or following disease progression on endocrine therapy. Patients may have also received up to one prior line of chemotherapy for advanced disease. Patients should have a good performance status, without active or uncontrolled metastases to the central nervous

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

system and can be of any menopausal status (Perimenopausal and premenopausal women must be treated with an LHRH agonist).

Clinical Notes:

- Treatment should continue until unacceptable toxicity or disease progression.
- Patients who progress ≤ 12 months from (neo) adjuvant therapy are eligible for treatment with ribociclib plus fulvestrant.
- Patients who experience disease progression on prior CDK 4/6 inhibitor therapy, fulvestrant or everolimus are not eligible for treatment with ribociclib with fulvestrant.
- Patients currently receiving fulvestrant monotherapy, and who have not progressed may have ribociclib added, provided they are CDK 4/6 inhibitor naïve and otherwise meet funding criteria.
- Patients who previously received everolimus plus exemestane will be eligible for funding of ribociclib plus fulvestrant on progression, provided that treatment was started prior to funding of CDK 4/6 + fulvestrant, patient must be CDK 4/6 naïve and otherwise meet funding criteria.
- Patients with prior adjuvant treatment with abemaciclib plus endocrine therapy are eligible if disease progression occurred 6 months or greater after completion of adjuvant abemaciclib.

RIFAXIMIN (*Zaxine 550mg Tablet*)

- For reducing the risk of overt hepatic encephalopathy (HE) recurrence if the following clinical criteria are met:
 - patients are unable to achieve adequate control of HE recurrence with lactulose alone;
 - used in combination with a maximal tolerated dose of lactulose.

RILUZOLE (*Rilutek 50mg Tablet and generic brands*)

- For the treatment of amyotrophic lateral sclerosis (ALS) or Lou Gehrig's Disease, when initiated by a neurologist with expertise in the management of ALS, when the patient has:
 - probable or definite diagnosis of ALS;
 - ALS symptoms for less than five years;
 - FVC $>60\%$ predicted upon initiation of therapy;
 - no tracheostomy for invasive ventilation.
- Coverage to be reviewed every six months.
- Coverage cannot be renewed once the patient has a tracheostomy for the purpose of invasive ventilation or mechanical ventilation.

RIOCIGUAT (*Adempas 0.5mg, 1.0mg, 1.5mg, 2.0mg, 2.5mg Tablet*)

- For the treatment of inoperable chronic thromboembolic pulmonary hypertension (CTEPH, World Health Organization [WHO] Group 4) or persistent or recurrent CTEPH after surgical treatment in adult patients (≥ 18 years of age) with WHO Functional Class (FC) II or III pulmonary hypertension (PH).
- Adempas should be prescribed by a clinician with experience in the diagnosis and treatment of CTEPH.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

***RIPRETINIB** (*Qinlock 50mg Tablet*)

- For the treatment of adult patients with advanced gastrointestinal stromal tumors (GIST) who have progression on or intolerance to imatinib, sunitinib and regorafenib.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should continue until disease progression or unacceptable toxicity.
- Patients must not have active CNS metastases.

Claim Note:

- Claims that exceed \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - Qinlock 50mg Tablet
 - 00900026
 - 00900027

RISANKIZUMAB (*Skyrizi 75mg/0.83mL Prefilled Injection, 150mg/mL Prefilled Syringe and 150mg/mL Prefilled Pen, 600mg/10ml Vial and 360mg/2.4ml Prefilled Cartridge Injection*)

PLAQUE PSORIASIS (PSO) (75MG/0.83ML PREFILLED INJECTION AND 150MG/ML PREFILLED SYRINGE/PEN)

- For patients with severe, debilitating chronic plaque psoriasis (PsO) who meet all of the following criteria:
 - Body Surface Area (BSA) involvement of >10% and/or significant involvement of the face, hands, feet or genitals
 - Failure to respond to, contraindication to or intolerant of methotrexate and cyclosporine
 - Failure to respond to, intolerant of or unable to access phototherapy
 - Written request of a dermatologist or prescriber with a specialty in dermatology
- Continued coverage is dependent on evidence of improvement, specifically:
 - $\geq 75\%$ reduction in the Psoriasis Area and Severity Index (PASI) score, OR
 - $\geq 50\%$ reduction in PASI with a ≥ 5 point improvement in DLQI (Dermatology Life Quality Index), OR
 - Significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals

Clinical Note:

- Treatment should be discontinued if a response has not been demonstrated by 16 weeks.

CROHN'S DISEASE (600MG/10ML VIAL AND 360MG/2.4ML PREFILLED CARTRIDGE INJECTION)

- For patients with moderate to severely active Crohn's disease and are refractory or have contraindications to an adequate course of 5-aminosalicylic acid and corticosteroids and other immunosuppressive therapy.

Clinical Note:

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

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Notes:

- Must be prescribed by a gastroenterologist or physician with a specialty in gastroenterology.
- Combined use of more than one biologic disease-modifying antirheumatic drugs (DMARD) will not be reimbursed.
- Initial reimbursement will be for intravenous doses of 600mg at Weeks 0, 4 and 8, with clinical response to be assessed prior to Week 12. Subsequent reimbursement for maintenance dosing is 360mg subcutaneously at Week 12, every 8 weeks thereafter.
- Initial Approval: 16 weeks
- Renewal Approval: 1 year

RISEDRONATE (*Actonel 30mg Tablet and generic brands*)

- Paget's disease of bone (2 month limit, one re-treatment course may be considered).
- Other requests reviewed on a case by case basis.

RISDIPLAM (*Evryssi 0.75mg/mL Powder for Solution*)**SPINAL MUSCULAR ATROPHY**

For patients diagnosed with 5q Spinal Muscular Atrophy (SMA) under the care of a specialist with experience in the diagnosis and management of SMA, if the following clinical criteria are met:

- Genetic documentation of 5q SMA homozygous gene deletion or compound heterozygote, AND
- Patients who:
 - are symptomatic and have genetic documentation of two or three copies of the SMN2 gene, AND
 - aged between 2 months and 7 months (inclusive), OR
 - aged 8 months up to 25 years and are non-ambulatory
- Patient is not currently requiring permanent invasive ventilation*, AND
- A baseline assessment using an age-appropriate scale (the Hammersmith Infant Neurological Examination [HINE] Section 2, Children's Hospital of Philadelphia Infant Test of Neuromuscular Disorders [CHOP INTEND], or Hammersmith Functional Motor Scale-Expanded [HFMSE]) must be completed prior to initiation of risdiplam treatment.
- For continued coverage, the patient must meet the following criteria:
 - There is demonstrated achievement or maintenance of motor milestone function (as assessed using age-appropriate scales: the [HINE] Section 2, CHOP INTEND, or HFMSE) after treatment initiation in patients aged between 2 months and 2 years at the time of treatment initiation; OR
 - There is demonstrated maintenance of motor milestone function (as assessed using age-appropriate scales: the HINE Section 2, CHOP INTEND, or HFMSE) after treatment initiation in patients aged between 2 years and 25 years at the time of treatment initiation; AND
 - Patient does not require permanent invasive ventilation*.

The decision to discontinue reimbursement should be based on 2 assessments separated by no longer than a 12-week interval.

Claim Notes:

- Coverage for risdiplam will not be provided in combination with other SMA drug therapies or post administration of onasemnogene abeparvovec.
- Approval: 12 months
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

transactions using the DIN first and then the following PINs:

- 00904768
- 00904769
- 00904770

* Permanent invasive ventilation is defined as the use of tracheostomy and a ventilator due to progression of SMA that is not due to an identifiable and reversible cause.

RISPERIDONE (*Risperdal Consta 12.5mg/2mL 25mg/2mL, 37.5mg/2mL, 50mg/2mL Injection*)

- For the treatment of patients who are:
 - not adherent to an oral antipsychotic, OR
 - currently receiving a long-acting injectable antipsychotic and require an alternative long-acting injectable antipsychotic.

Claim Note:

- Requests will not be considered for the treatment of psychotic symptoms related to dementia.

RITUXIMAB (*Riximyo, Ruxience & Truxima 10mg/mL Injection*)

- For the treatment of rheumatoid arthritis¹, vasculitis², or other autoimmune diseases³.

Clinical Notes:

1. Severe intolerance or other contraindication to an anti-TNF agent or failed an adequate trial of an anti-TNF agent.
2. Severe intolerance or other contraindication to cyclophosphamide or failed an adequate trial of cyclophosphamide.
3. Previously failed treatments must be provided if applicable.

Claims Notes:

- Must be prescribed by a specialist.
- Approval period: long term

ROMOSOZUMAB (*Evenity 90mg/mL Prefilled Syringe*)

- For the treatment of osteoporosis in postmenopausal women who meet all the following criteria:
 - Have a history of osteoporotic fracture; and
 - Are at high risk for future fracture, defined as a 10-year fracture risk $\geq 20\%$ as per the Fracture Risk Assessment (FRAX) tool; and
 - Are treatment naive to osteoporosis medications, except for calcium and/or vitamin D.

Claim Notes:

- Maximum approval period: 12 months per lifetime.
- Concurrent use with other osteoporosis medications, except for calcium and/or vitamin D, will not be reimbursed.

ROTIGOTINE (*Neupro 2mg/24hr, 4mg/24hr, 6mg/24hr, 8mg/24hr Patch*)

- For adjunctive therapy to levodopa for the treatment of patients with advanced stage Parkinson's disease (APD).

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

RUFINAMIDE (*Banzel 100mg, 200mg and 400mg Tablet*)

- For the adjunctive treatment of seizures associated with Lennox-Gastaut syndrome for patients who meet all of the following criteria:
 - are under the care of a physician experienced in treating Lennox-Gastaut syndrome-associated seizures, AND
 - are currently receiving two or more antiepileptic drugs, AND
 - in whom less costly antiepileptic drugs are ineffective or not appropriate.

***RUXOLITINIB** (*Jakavi 5mg, 10mg, 15mg and 20mg Tablet*)

ACUTE GRAFT-VERSUS-HOST DISEASE

- For the treatment of steroid-refractory or steroid-dependent acute graft-versus-host disease (aGvHD) in adult and pediatric patients aged 12 years and older who meet all the following criteria:
 - Clinically diagnosed grade II to IV aGvHD according to the NIH criteria (Harris et al. [2016]).
 - Confirmed diagnosis of corticosteroid-refractory or corticosteroid-dependent aGvHD.

Renewal criteria:

- Achieved an overall response (i.e., CR, VGPR, PR, or stable disease with significant reduction in steroid doses), according to standard NIH criteria at day 28.
- For subsequent renewals, patients should be assessed for treatment response every 2 to 3 months, until the occurrence of any of the discontinuation criteria listed below.

Clinical Notes:

- Treatment should be discontinued upon the occurrence of any of the following:
 - progression of aGvHD, defined as worsening of aGvHD symptoms or occurrence of new aGvHD symptoms
 - unacceptable toxicity
 - addition of systemic therapies (other than calcineurin inhibitors) for aGvHD after day 28
 - recurrence or relapse of underlying hematological malignancy.

Claim Notes:

- Must be prescribed by clinicians who have experience in the diagnosis and management of patients with aGvHD.
- Must not be added to patients' concurrent treatment of systemic therapies for the treatment of aGvHD other than steroids with or without calcineurin inhibitors.
- Approval: 6 months

CHRONIC GRAFT-VERSUS-HOST DISEASE

- For the treatment of chronic graft-versus-host disease (cGvHD) in adults and pediatric patients aged 12 years and older who have inadequate response to corticosteroids or other systemic therapies who meet all the following criteria:
 - Clinically diagnosed cGvHD staging of moderate to severe based on NIH consensus criteria
 - Confirmed diagnosis cGvHD with inadequate response to corticosteroids or other systemic therapies

Renewal criteria:

- Achieved an overall response (i.e., CR or PR, or stable disease with significant reduction in steroid doses), according to NIH criteria, after 24 weeks of therapy

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

- Treatment should be discontinued upon the occurrence of any of the following:
 - Progression of cGvHD, defined as worsening of cGvHD symptoms or occurrence of new cGvHD symptoms
 - recurrence or relapse of underlying hematological malignancy

Claim Notes:

- Must be prescribed by clinicians who have experience in the diagnosis and management of patients with cGvHD.
- Must not be added to patients' concurrent treatment of systemic therapies other than steroids with or without calcineurin inhibitors.
- Initial Approval: 6 months
- Renewal Approval: 1 year

MYELOFIBROSIS

- As a single agent in patients with intermediate or high risk symptomatic myelofibrosis (using the Dynamic International Prognostic Scoring System (DIPSS) Plus or symptomatic splenomegaly) with an ECOG performance status (PS) ≤ 3 as **first line therapy or refractory to other treatments**. Ongoing monitoring and follow up of therapy will be required.

POLYCYTHEMIA VERA

- For the treatment of patients with polycythemia vera who have demonstrated resistance or intolerance to hydroxyurea (HU).

Renewal Criteria:

- Written confirmation that the patient has responded to treatment and there is no evidence of disease progression.

Clinical Notes:

1. Patients must have a good performance status.
2. Treatment should be discontinued upon disease progression or unacceptable toxicity.
3. Resistance is considered if, after at least 3 months of HU therapy at the maximum tolerated dose, patients experience at least one of the following:
 - Need for phlebotomy to maintain hematocrit (HCT) $< 45\%$
 - Uncontrolled myeloproliferation (i.e., platelet count $> 400 \times 10^9/L$ and white blood cell count $> 10 \times 10^9/L$)
 - Failure to reduce massive splenomegaly by greater than 50%, as measured by palpation
4. Intolerance to HU is considered if patients experience at least one of the following:
 - Absolute neutrophil count $< 1.0 \times 10^9/L$, platelet count $< 100 \times 10^9/L$ or hemoglobin $< 100g/L$ at the lowest dose of HU required to achieve a response (a response to HU is defined as HCT $< 45\%$ without phlebotomy, and/or all of the following: platelet count $< 400 \times 10^9/L$, white blood cell count $< 10 \times 10^9/L$, and nonpalpable spleen).
 - Presence of leg ulcers or other unacceptable HU-related non-hematological toxicities (defined as grade 3 or 4 or, more than one week of grade 2) such as mucocutaneous manifestations, gastrointestinal symptoms, pneumonitis, or fever.
 - Toxicity requiring permanent discontinuation of HU, interruption of HU until toxicity resolved, or hospitalization due to HU toxicity.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Notes:

- Initial approval period: 6 months
- Renewal approval period: 1 year

***SACCHARATED IRON OXIDE** (*Venofer 20mg/mL Injection and generic brands*)

- For the treatment of iron deficiency anemia in patients intolerant to oral iron replacement products; OR
- For patients who have not responded to adequate therapy with oral iron.

Notes:

- Given the safety concerns associated with IV iron, it is expected that the patients will be carefully screened and will have tried various oral iron options before being eligible for IV iron.
- Details regarding oral iron tried, length of therapy, and outcome must be provided.

SACUBITRIL/VALSARTAN (*Entresto 24.3mg/25.7mg, 48.6mg/51.4mg, 97.2mg/102.8mg Tablet*)

- For the treatment of heart failure (HF) with reduced ejection fraction in patients with New York Heart Association (NYHA) class II or III HF to reduce the incidence of cardiovascular (CV) death and HF hospitalization, if ALL of the following clinical criteria are met:
 - Reduced left ventricular ejection fraction (LVEF) (< 40%);
 - Patient has NYHA class II to III symptoms despite at least four weeks of treatment with stable doses of all of the following medications:
 - an angiotensin-converting enzyme inhibitor (ACEI) or an angiotensin II receptor antagonist (ARB);
 - a beta blocker;
 - other recommended therapies, including an aldosterone antagonist (if tolerable);
- Plasma B-type natriuretic peptide (BNP) ≥ 150 pg/mL or N-terminal prohormone B-type natriuretic peptide (NT-proBNP) ≥ 600 pg/mL; or plasma BNP ≥ 100 pg/mL or NT-proBNP ≥ 400 pg/mL levels if the patient has been hospitalized for HF within the past 12 months. If BNP testing is not accessible the reasons must be clearly outlined.

Clinical Notes:

- Initiation and up-titration should be conducted by a prescriber experienced with the treatment of heart failure
- For patients who have not received four weeks of therapy with a beta blocker or aldosterone antagonist due to an intolerance or contraindication, details must be provided.

***SALBUTAMOL** (*0.5mg/mL, 1mg/mL, 2mg/mL Unit Dose Inhaled Solution and 5mg/mL Inhaled Solution*)

- See [Wet Nebulization Solutions](#)

***SALBUTAMOL, IN COMBINATION** (*Combivent Inhaled Solution and generic brands*)

- See [Wet Nebulization Solutions](#)

SALMETEROL (*Serevent 50mcg/dose Diskus*)

- See [Long-Acting Beta₂-Agonists](#)

SALMETEROL IN COMBINATION (*Advair 50/100mcg, 50/250mcg, 50/500mcg Diskus, generic brands and HFA 25/125 mcg/dose, HFA 25/250 mcg/dose Inhaler*)

- See [Long-Acting Beta₂-Agonists/Inhaled Corticosteroids](#)

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

SAPROPTERIN DIHYDROCHLORIDE (*Kuvan 100mg Tablet and 100mg and 500mg Sachet*)

- Ongoing funding of Kuvan will be considered for nonpregnant patients and patients actively planning pregnancy who have a diagnosis of Phenylketonuria (PKU) and who have demonstrated a response to the initial 6 month trial of sapropterin [reimbursed through the Supplier's Patient Support Program (PSP) 'BioMarin RareConnections']

Inclusion Criteria for entry into the 6 month trial period:

- For the management of patients with the diagnosis of hyperphenylalaninemia (HPA) due to tetrahydrobiopterin (BH4)-responsive phenylketonuria (PKU) who meet ALL of the following criteria:
 - A diagnosis of Phenylketonuria (PKU) confirmed through an approved test.
 - Compliance with a low protein diet and formulas.
 - Baseline blood phenylalanine (Phe) levels > 360 µmol/L despite compliance with a low protein diet (require at least 2 baseline levels during a 3 to 6 month time frame).
 - Baseline protein intake assessment by a dietitian.
 - Ability to comply with medication regimen.
 - Managed by a physician specialized in metabolic/biochemical diseases.
- Modified Criteria for Pregnant Patients during the 6 month trial period:
 - Patient has a diagnosis of PKU confirmed through an approved test
 - Patient's treatment is being managed by a prescriber specialized in metabolic/biochemical diseases; and
 - Patient's baseline blood Phe level is greater than 360 µmol/L despite compliance with all recommendations for dietary intervention and monitoring or compliance with a low protein diet.
- Patients will be eligible for funding through the Nova Scotia Pharmacare Programs after demonstrating a response to the 6 month trial period, as per the trial criteria.

Initial Criteria Post 6 Month Trial:

- For the management of patients with the diagnosis of hyperphenylalaninemia (HPA) due to tetrahydrobiopterin (BH4)-responsive phenylketonuria (PKU) who meet ALL of the following criteria:
 - Compliance with low protein diet, formulas, and Kuvan; AND
 - During the 6 month trial period under the patient support program BioMarin RareConnections, patient has achieved a demonstrated response to the Kuvan responsiveness test or PKU clinical protocol, based on the following information:
 - the clinic's definition for response; and
 - all relevant laboratory results used to determine that the Patient was a responder to Kuvan
 - Patient meets one of the following:
 - normal sustained Blood Phe levels [> 120 µmol/L and < 360 µmol/L] (At least 2 levels measured at least 1 month apart); OR
 - sustained blood Phe reduction of at least 30% (At least 2 levels measured at least 1 month apart) compared to baseline if the Phe baseline level is < 1200 µmol/L; OR
 - sustained blood Phe reduction of at least 50% (At least 2 levels measured at least 1 month apart) compared to baseline if the Phe baseline level is > 1200 µmol/L;
 - Demonstrated an increase in dietary protein tolerance based on targets set between the clinician and patient
 - Managed by a prescriber specialized in metabolic/ biochemical diseases.
- Dosage: Up to a maximum of 20 mg/kg per day
- Approval Duration: 1 year

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Renewal Criteria:

- Renewals will be considered for patients who demonstrate ongoing response to treatment.
- Renewal Approval Duration: 1 year

Claim Note:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - Kuvan 100mg Tablet 00904580
 - Kuvan 100mg Sachet 00904805
 - Kuvan 500mg Sachet 00904579

SATRALIZUMAB (*Enspryng 120mg/mL Prefilled Syringe*)**NEUROMYELITIS OPTICA SPECTRUM DISORDER**

- For the treatment of patients 12 years of age and older with neuromyelitis optica spectrum disorder (NMOSD) who meet all of the following criteria:
 - Are anti-aquaporin 4 (AQP4) seropositive
 - Must have had at least one relapse of NMOSD in the previous 12 months:
 - despite an adequate trial of other accessible preventive treatments¹ for NMOSD, OR
 - because the patient cannot tolerate other preventive treatments¹ for NMOSD
- Patients must have an EDSS score of 6.5 points or less.
- Satralizumab should not be initiated during a NMOSD relapse episode.

Renewal:

- Requests for renewal will be considered for patients who maintain an EDSS score of less than 8 points.

Clinical Note:

- Must be prescribed by a neurologist with expertise in treating NMOSD.

Claim Notes:

- Approval Period: 1 year
- Combined use of more than one biologic drug will not be reimbursed.
- Approvals will be for a maximum of 120mg at week 0, 2 and 4, then 120 mg every four weeks thereafter.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00904802

¹Other accessible preventative treatments include, but are not limited to, monoclonal antibodies including rituximab and other immunosuppressants.

SARILUMAB (*Kevzara 150mg/1.14mL, 200mg/1.14mL Prefilled Pen & 150mg/1.14mL, 200mg/1.14mL Prefilled Syringe*)

- For the treatment of severely active rheumatoid arthritis, in combination with methotrexate or other disease modifying antirheumatic drugs (DMARDs), in adult patients who are refractory or intolerant to:
 - methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age), OR

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- use in combination with another DMARD, for a minimum of 12 weeks; AND
- methotrexate in combination with at least two other DMARDs, such as hydroxychloroquine and sulfasalazine, for a minimum of 12 weeks.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- Optimal treatment response to DMARDs may take up to 24 weeks, however coverage of a biologic therapy can be considered if no improvement is seen after 12 weeks of triple DMARD use.
- If patient factors (e.g. intolerance) prevent the use of triple DMARD therapy, these must be described and dual therapy with DMARDs must be tried.
- 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. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Combined use of more than one biologic DMARD will not be reimbursed.
- Initial Approval: 6 months.
- Renewal Approval: 1 year. Confirmation of continued response is required.

SAXAGLIPTIN (*Onglyza 2.5mg and 5mg Tablet and generic brands*)

- For the treatment of Type II diabetes for patients with:
 - inadequate glycemic control on metformin and a sulfonylurea; and
 - for whom insulin is not an option.

SAXAGLIPTIN AND METFORMIN (*Komboglyze 2.5mg/500mg, 2.5mg/850mg, 2.5mg/1000mg Tablet*)

- For the treatment of Type II diabetes for patients:
 - who are already stabilized on therapy with metformin, a sulfonylurea and saxagliptin to replace the individual components of saxagliptin and metformin; and
 - for whom insulin is not an option.

SEBELIPASE ALFA (*Kanuma 2mg/mL IV Solution*)

- For the treatment of patients diagnosed with lysosomal acid lipase (LAL) deficiency who meet all of the following criteria:
 - Documented biochemical evidence of deficient LAL activity and two documented pathogenic mutations in the LIPA gene
 - AND**
 - Patients who:
 - Have onset of clinical manifestations¹ of LAL deficiency before six months of age.
 - OR**
 - Have at least one of the following clinical manifestations¹ of LAL deficiency at 6 months of age and older:
 - Persistently elevated transaminases (ALT > 1.5 x ULN² or AST > 1.5 x ULN²) as

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

measured by two assessments three to six months apart.

- Persistent dyslipidemia (LDL-c and/or TG values in the top 5th percentile based on sex and age) as measured by two assessments three to six months apart.
- Any documented hepatomegaly or hepatosplenomegaly.
- Liver fibrosis confirmed by biopsy.
- Failure to thrive.
- Growth impairment³.
- Evidence of intestinal affection and/or malabsorption.

AND

- Must not demonstrate evidence of any of the following:
 - Increased portal vein pressures, or de novo evidence of portal hypertension on ultrasound and Doppler, or new clinical presentation of portal hypertension (e.g., esophageal varices).
 - Severe hepatic dysfunction (Child-Pugh Class C).
 - End-stage liver disease.

Discontinuation Criteria:

- For patients with onset of clinical manifestations of LAL deficiency at six months of age and older if the patient:
 - Progresses to end-stage liver failure or multi-organ failure.

OR

- Has at least three out of the following response components compared to baseline values after 12 months of therapy:
 - Less than 10% improvement in ALT or AST.
 - Worsening of liver fibrosis confirmed by biopsy.
 - Persisting growth impairment³ despite sebelipase alfa therapy and nutritional interventions.
 - At least a 15% increase in spleen volume and/or a greater than 15% increase in liver volume on ultrasound.
 - Increased portal vein pressures, or de novo evidence of portal hypertension on ultrasound and Doppler, or new clinical presentation of portal hypertension (e.g., esophageal varices).
- Regardless of age of onset, for adverse events from sebelipase alfa (particularly hypersensitivity reactions including anaphylaxis, hypotension, or fever), which cannot be managed with standard treatment and/or have a significant impact on the patient's quality of life or are life-threatening.

Clinical Notes:

1. The physician must provide baseline values for the clinical manifestation at the time of initial request for reimbursement.
2. Based on age- and- sex-specific normal values for ALT and AST.
3. Growth impairment is defined as decreased body weight across at least two of the major centiles on a WHO weight-for-age chart, or body weight below 10th centile and no weight gain within two weeks and/or decreased height across at least two of the major centiles on a WHO height-for-age chart.

Claim Notes:

- The patient must be under the care of a specialist with experience in the diagnosis and management of LAL deficiency.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Initial Approval: 12 months.
- Renewals: 6 months.
- Claims for Kanuma 2mg/mL IV Solution that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904599
 - 00904600
 - 00904601

Please call the Nova Scotia Pharmacare Programs if additional PINs are required.

SECUKINUMAB (*Cosentyx 300mg dose kits (two subcutaneous injections of 150mg/1mL)*)

ANKYLOSING SPONDYLITIS

- For the treatment of patients with moderate to severe ankylosing spondylitis (Bath AS Disease Activity Index (BASDAI) score ≥ 4 on 10 point scale) who:
 - Have axial symptoms¹ and who have failed to respond to the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each, or in whom NSAIDs are contraindicated;
 - OR
 - Have peripheral symptoms and who have failed to respond to, or have contraindications to, the sequential use of at least 2 NSAIDs at the optimum dose for a minimum period of 2 weeks each and have had an inadequate response to an optimal dose or maximal tolerated dose of a DMARD.
- Requests for renewal must include information demonstrating the beneficial effects of the treatment, specifically:
 - A decrease of at least 2 points on the BASDAI scale, compared with the pre-treatment score, or
 - Patient and expert opinion of an adequate clinical response as indicated by a significant functional improvement (measured by outcomes such as HAQ or “ability to return to work”).

Clinical Note:

- Patients with recurrent uveitis (2 or more episodes within 12 months) as a complication to axial disease do not require a trial of NSAIDs alone.

Claim Notes:

- Must be prescribed by a rheumatologist or prescriber with a specialty in rheumatology.
- Combined use of more than one biologic DMARD will not be reimbursed.
- Approvals will be for 150mg given at weeks 0, 1, 2, 3, and 4, followed by monthly maintenance dosing. If a patient continues to have active ankylosing spondylitis, a monthly maintenance dosage of 300 mg may be considered.
- Each 300 mg dose is given as two subcutaneous injections of 150 mg.
- Initial Approval: 6 months.
- Renewal Approval: 1 year.

PSORIASIS

- For patients with severe, debilitating chronic plaque psoriasis who meet all of the following:
 - Body surface area (BSA) involvement of $>10\%$ and/or significant involvement of the face, hands, feet or genitals;
 - Failure to, contraindication to or intolerant of methotrexate and cyclosporine;

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Failure to, intolerant of or unable to access phototherapy;
- Written request of a dermatologist or prescriber with a specialty in dermatology.
- Continued coverage is dependent on evidence of improvement, specifically:
 - A >75% reduction in the Psoriasis Area and Severity Index (PASI) score; or
 - A >50% reduction in PASI with a > 5 point improvement in DLQI (Dermatology Life Quality Index); or
 - Significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals.
- Concurrent use of biologics not approved.
- Initial approval for a maximum of 12 weeks.
- Coverage may be approved as follows: initial dosing of 300 mg doses at Weeks 0, 1, 2 and 3, followed by monthly maintenance dosing of 300 mg doses starting at Week 4.

PSORIATIC ARTHRITIS

- For the treatment of patients with predominantly axial psoriatic arthritis who are refractory, intolerant or have contraindications to the sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each.
- For the treatment of patients with predominantly peripheral psoriatic arthritis who are refractory, intolerant or have contraindications to:
 - The sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each; and
 - Methotrexate (oral or parenteral) at a dose of $\geq 20\text{mg}$ weekly ($\geq 15\text{mg}$ if patient is ≥ 65 years of age) for a minimum of 8 weeks; and
 - Leflunomide for a minimum of 10 weeks or sulfasalazine for a minimum of 3 months

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- 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 to treatments. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Combined use of more than one biologic DMARD will not be reimbursed.
- Approvals will be for a maximum of 150mg given at weeks 0, 1, 2, 3, and 4, then monthly. Requests for 300mg monthly will be considered for patients who have previously had an inadequate response to TNF-inhibitors.
- Initial approval: 6 months.
- Renewal Approval: 1 year. Confirmation of continued response is required.

SELECTIVE 5HT₁ - RECEPTOR AGONISTS (*Almotriptan Tablet, Naratriptan Tablet, Sumatriptan Nasal Spray*)

SUMATRIPTAN 50MG & 100MG TABLET, NARATRIPTAN TABLET

- For the treatment of migraine¹ headache when:
 - migraines are moderate² in severity and other therapies (e.g. NSAIDs, acetaminophen, DHE spray) are

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

not effective; or

- migraine attacks are severe² or ultra severe².

ALMOTRIPTAN TABLET, ZOLMITRIPTAN NASAL SPRAY, SUMATRIPTAN NASAL SPRAY

- For the treatment of migraine¹ headache of moderate² intensity when:
 - Other therapies (e.g. NSAIDs, acetaminophen, DHE spray) are not effective AND patients have not responded to oral sumatriptan, zolmitriptan, rizatriptan and naratriptan.
 - For the treatment of migraine¹ headache of severe² or ultra severe² intensity when patients have not responded to oral sumatriptan, zolmitriptan, rizatriptan, and/or naratriptan.

SUMATRIPTAN 6MG/SYRINGE INJECTION

- For the treatment of migraine¹ headache of moderate² intensity when:
 - other therapies (e.g. NSAIDs, acetaminophen, DHE spray) are not effective AND oral and nasal triptans are not appropriate.
 - for the treatment of migraine¹ headache of severe² or ultra severe² intensity when oral and nasal triptans are not appropriate.

Notes:

- Coverage limited to 18 doses/3 months³
 - patients with >3 migraines/month on average despite prophylactic therapy may be considered for up to a maximum of 12 doses/30 day
1. As diagnosed based on current Canadian guidelines.
 2. Definitions: Moderate – pain is distracting causing need to slow down and limit activities;
Severe – pain affects ability to concentrate and very difficult to continue with daily activities;
Ultra severe – unable to speak or think clearly; not able to function; likely lying down or sleeping.
 3. Reimbursement will be available for a maximum quantity of 18 triptan doses per quarter (e.g., Jan to Mar) regardless of the agent(s) used within the 90 day period.

SELEXIPAG (*Uptravi 200mcg, 400mcg, 600mcg, 800mcg, 1000mcg, 1200mcg, 1400mcg, 1600mcg Tablet*)

- For the long-term treatment of idiopathic pulmonary arterial hypertension (PAH), heritable HPAH, PAH associated with connective tissue disorders, and PAH associated with congenital heart disease, in adult patients with World Health Organization (WHO) functional class (FC) II to III to delay disease progression, if the following clinical criteria are met:
 - Inadequate control with a first- and second-line PAH therapy.
 - Must be prescribed by a clinician with experience in the diagnosis and treatment of PAH.

Claim Note:

- Combination therapy with prostacyclin or prostacyclin analogs will not be reimbursed.

***SELINEXOR (*Xpovio 20mg Tablet*)**

- In combination with bortezomib and dexamethasone for the treatment of adult patients with multiple myeloma and who have received at least one prior therapy.

Clinical Notes:

1. Prior treatment with bortezomib/proteasome inhibitor is permitted if all the following criteria are met:
 - Best response achieved with bortezomib/proteasome inhibitor was at least a partial response
 - Bortezomib/proteasome inhibitor not discontinued for grade 3 or higher toxicity
 - Bortezomib/proteasome inhibitor treatment-free interval has been at least six months.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

2. Treatment should continue until disease progression or unacceptable toxicity.

Claim Note:

- Claims that exceed \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00900031

SELPERCATINIB (*Retevmo 40mg and 80mg Capsule*)*MEDULLARY THYROID CANCER**

- For the treatment of patients 12 years and older with unresectable locally advanced or metastatic RET-mutant medullary thyroid cancer (MTC) who have progressed on, are intolerant to, or have a contraindication to first-line therapy.

Clinical Notes:

- Discontinuation for unacceptable toxicity or loss of clinical benefit.
- Patients should have a good performance status.
- Monotherapy only.
- Confirm RET mutation prior to initiating therapy.
- Patients with prior progression on a RET inhibitor are ineligible.

DIFFERENTIATED THYROID CARCINOMA

- For the treatment of adult patients with locally advanced or metastatic RET fusion-positive differentiated thyroid carcinoma (DTC) not amenable to surgery or radioactive iodine therapy following prior Lenvatinib and/or Sorafenib.

Clinical Notes:

- Discontinuation for unacceptable toxicity or loss of clinical benefit.
- Patients should have a good performance status.
- Monotherapy only.
- Confirm RET mutation prior to initiating therapy.
- Patients with prior progression on a RET inhibitor are ineligible.

NON-SMALL CELL LUNG CANCER

- For the treatment of adult patients with metastatic RET fusion-positive non-small cell lung cancer (NSCLC) as first-line treatment or after prior systemic therapy.

Clinical Notes:

- Discontinuation for unacceptable toxicity or loss of clinical benefit.
- Patients should have a good performance status.
- Monotherapy only.
- Confirm RET mutation prior to initiating therapy.
- Patients with prior progression on a RET inhibitor are ineligible.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Note:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - Retevmo 80mg Cap
 - 00904987

SEMAGLUTIDE (Ozempic 2mg/1.5mL, 2mg/3mL and 4mg/3mL Prefilled Pen)

- For the treatment of type 2 diabetes in combination with metformin and a sulfonylurea, when diet and exercise plus dual therapy with metformin and a sulfonylurea do not achieve adequate glycemic control.

Claim Note:

- Approvals will be for a maximum of 1 prefilled pen every 4 weeks.

SEMAGLUTIDE (Rybelsus 3mg, 7mg and 14mg Tablet)

- For the treatment of type 2 diabetes in combination with metformin and a sulfonylurea, when diet and exercise plus dual therapy with metformin and a sulfonylurea do not achieve adequate glycemic control.

Claim Note:

- Approvals will be for a maximum of one tablet per day.

SEROTONIN (5-HT₃) ANTAGONISTS (Granisetron, Ondansetron)*EMESIS (GRANISETRON, ONDANSETRON)**

- For the treatment of emesis in patients who are:
 - receiving moderately or severely emetogenic chemotherapy [**Criteria Code 01**] or
 - receiving intravenous chemotherapy or radiotherapy and who have not experienced adequate control with other available antiemetics [**Criteria Code 02**] or
 - receiving intravenous chemotherapy or radiotherapy and who are experiencing intolerable side effects to other antiemetics, including steroids and anti-dopaminergic agents [**Criteria Code 03**]

Note:

- Use of criteria codes is limited to appropriate doses pre and post chemotherapy or radiation. Criteria codes must not be used for claims related to other causes of nausea and vomiting or for long term, daily management of nausea and vomiting.

NAUSEA, VOMITING IN PEDIATRIC PATIENTS (ONDANSETRON)

- For the treatment of nausea and vomiting in pediatric patients (under 18 years of age) receiving chemotherapy (e.g., methotrexate) for chronic non-oncology conditions who have experienced an episode of nausea/emesis. [**Criteria Code 04**]

SEVELAMER (800mg Tablet and generic brands)

- For the treatment of hyperphosphatemia (>1.8 mmol/L) in patients with end-stage renal disease (eGFR < 15 mL/min) who have:
 - Inadequate control of phosphate levels on a calcium based phosphate binder, or
 - Hypercalcemia (corrected for albumin), or
 - Calciphylaxis (calcific arteriopathy)

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Notes:

- Must be prescribed by a nephrologist or other prescriber within the Provincial Dialysis Program.
- Initial Approval: 6 months.
- Renewal Approval: 1 year. Confirmation of improvement of phosphate levels is required (lab values must be provided).

SILDENAFIL (*Revatio 20mg Tablet and generic brands*)

- Written request from a pulmonary arterial hypertension (PAH) specialist only
- Diagnosis of PAH should be confirmed by right heart catheterization
- Dose of sildenafil will be limited to 20mg tid

IPAH (FUNCTIONAL CLASS III)

- For the treatment of patients with World Health Organization (WHO) functional class III idiopathic pulmonary arterial hypertension (IPAH) who do not demonstrate vasoreactivity on testing, or who do demonstrate vasoreactivity on testing but fail a trial of calcium channel blockers (CCB), or are intolerant to CCB.

PAH SECONDARY TO CONNECTIVE TISSUE DISEASE (FUNCTIONAL CLASS III)

- For the treatment of patients with World Health Organization (WHO) functional class III pulmonary hypertension associated with connective tissue disease who do not respond to conventional therapy.

SIPONIMOD (*Mayzent 0.25mg and 2mg Tab*)**SECONDARY PROGRESSIVE MULTIPLE SCLEROSIS****Initiation Criteria:**

- For the treatment of patients with active secondary progressive multiple sclerosis, who meet all the following criteria:
 - a history of relapsing-remitting multiple sclerosis (RRMS)
 - an Expanded Disability Status Scale (EDSS) score of 3.0 to 6.5
 - documented EDSS progression during the two years prior to initiating treatment with siponimod (≥ 1 point if EDSS < 6.0; ≥ 0.5 points if EDSS ≥ 6.0 at screening).

Renewal Criteria:

- Patients who do NOT exhibit evidence of disease progression since the previous assessment. Disease progression is defined as:
 - an increase in the EDSS score of greater than or equal to 1 point if the EDSS score was 3.0 to 5.0 at siponimod initiation
 - OR
 - an increase of greater than or equal to 0.5 points if the EDSS score was 5.5 to 6.5 at siponimod initiation
- Patients who do NOT exhibit one of the following:
 - progression to an EDSS score of equal to or greater than 7.0 at any time during siponimod treatment
 - confirmed worsening of at least 20% on the timed 25-foot walk (T25W) since initiating siponimod treatment

Clinical Note:

- Patients should be assessed for a response to siponimod every six months.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claims Notes:

- The patient is under the care of a neurologist with experience in the diagnosis and management of multiple sclerosis.
- Siponimod should not be used in combination with other disease-modifying treatments (DMTs) used to treat multiple sclerosis.
- Approval period: 1 year

SITAGLIPTIN (*Januvia 25mg, 50mg, 100mg Tablet and generic brands*)

- For the treatment of Type II diabetes for patients with:
 - inadequate glycemic control on metformin and a sulfonylurea; and
 - for whom insulin is not an option.

SITAGLIPTIN AND METFORMIN (*Janumet 50/500mg, 50/850mg, 50/1000mg, 50mg/500mg XR, 50/1000 XR and 100mg/1000mg XR Tablet and generic brands*)

- For the treatment of Type II diabetes for patients:
 - who are already stabilized on therapy with metformin, a sulfonylurea and sitagliptin to replace the individual components of sitagliptin and metformin;
AND
 - for whom insulin is not an option.

SMOKING CESSATION THERAPIES (*bupropion and varenicline*)

- A maximum of 12 weeks standard therapy (168 tablets) will be reimbursed annually without a special authorization request.
- Additional reimbursement (e.g. for a second course of therapy) will require a special authorization request with details regarding readiness to quit, success with previous therapy, enrolment in cessation programs and any other pertinent information.

***SODIUM FERRIC GLUCONATE** (*Ferrlecit 12.5mg/mL Injection*)

- For the treatment of iron deficiency anemia in patients intolerant to oral iron replacement products; OR
- For patients who have not responded to adequate therapy with oral iron.

Notes:

- Given the safety concerns associated with IV iron, it is expected that the patients will be carefully screened and will have tried various oral iron options before being eligible for IV iron.
- Details regarding oral iron tried, length of therapy, and outcome must be provided.

SODIUM PHENYLBUTYRATE (*Pheburane 483mg/g Oral Granules*)

- For the treatment of patients with urea cycle disorders (UCDs).

Clinical Note:

- Diagnosis must be confirmed by blood, enzymatic, biochemical or genetic testing.

Claim Note:

- Must be prescribed by, or in consultation with, a physician experienced in the treatment of UCDs.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

SODIUM PHENYL BUTYRATE AND URSODOXICOLTAURINE (*Albrioza 3g/1g Sachet*)

- For the treatment of amyotrophic lateral sclerosis (ALS), if the following criteria are met:

Initiation:

- Patient with a diagnosis of definite ALS; AND
- Patient who meets all of the following:
 - have had ALS symptoms for 18 months or less
 - have a forced vital capacity of at least 60% of predicted value
 - not require permanent non-invasive ventilation or invasive ventilation

Renewal:

- Reimbursement of treatment should be discontinued in patients who meet any one of the following criteria:
 - the patient becomes non-ambulatory and is unable to cut food and feed themselves without assistance, irrespective of whether a gastrostomy is in place; OR
 - patient requires permanent non-invasive ventilation

Claim Notes:

- Patient must be under the care of a specialist with experience in the diagnosis and management of ALS.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00904825

SOFOSBUVIR (*Sovaldi 400mg Tablet*)

- For treatment-naïve or treatment-experienced adult patients with chronic hepatitis C virus (HCV) who meet the following criteria:

Approval Period and Regimen

Genotype 2

- | | |
|------------------------------|--|
| ▪ Without cirrhosis | ▪ 12 weeks in combination with ribavirin (RBV) |
| ▪ With compensated cirrhosis | |

Genotype 3

- | | |
|------------------------------|------------------------------------|
| ▪ Without cirrhosis | ▪ 24 weeks in combination with RBV |
| ▪ With compensated cirrhosis | |

Genotype 3

- | | |
|---------------------|--|
| ▪ Without cirrhosis | ▪ 12 weeks in combination with daclatasvir |
|---------------------|--|

Genotype 3

- | | |
|---|--|
| ▪ With compensated or decompensated cirrhosis | ▪ 12 weeks in combination with daclatasvir and RBV |
| ▪ Post-liver transplant without cirrhosis or with compensated cirrhosis | |

- **Patients must also meet all of the following criteria:**

- Must be prescribed by a hepatologist, gastroenterologist, or infectious disease specialist (or other physician experienced in treating a patient with hepatitis C infection)
- Lab-confirmed hepatitis C genotype 2 and 3
- Quantitative HCV RNA value within the last 6 months
- Fibrosis stage must be provided

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

1. Treatment-experienced is defined as a patient who has been previously treated with a peginterferon/ribavirin regimen and has not experienced an adequate response.
2. Acceptable methods for the measurement of fibrosis score include Fibrotest, liver biopsy, transient elastography (FibroScan®), serum biomarker panels (such as AST-to-Platelet Ratio Index or Fibrosis-4 score) either alone or in combination.
3. Compensated cirrhosis is defined as a CTP score of 5 to 6 (Class A) and decompensated cirrhosis as a CTP score of 7 or above (Class B or C).
4. Re-treatment for direct-acting antiviral failures will be considered on a case-by-case basis.

Claim Notes:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904041
 - 00904042
- Claims will be limited to a 28-day supply.

** [Criteria Code 34] has been added for use effective December 1, 2019 to allow payment of a patient's initial 28 day supply only. Criteria code 34 should be provided by the prescribing physician only.

SOFOSBUVIR AND VELPATASVIR (*Epclusa 400mg/100mg Tablet*)

- For treatment-naïve or treatment-experienced adult patients with chronic hepatitis C virus (HCV) who meet the following criteria:

Approval Period and Regimen**Genotypes 1, 2, 3, 4, 5, 6 or mixed genotypes**

- | | |
|---------------------------------------|------------|
| ▪ Patients with compensated cirrhosis | ▪ 12 weeks |
| ▪ Patients without cirrhosis | |

Genotypes 1, 2, 3, 4, 5, 6 or mixed genotypes

- | | |
|---|--|
| ▪ Patients with decompensated cirrhosis | ▪ 12 weeks in combination with ribavirin |
|---|--|
- **Patients must also meet all of the following criteria:**
 - Must be prescribed by a hepatologist, gastroenterologist, or infectious disease specialist (or other physician experienced in treating a patient with hepatitis C infection)
 - Lab-confirmed hepatitis C genotype 1, 2, 3, 4, 5, 6 or mixed genotypes
 - Quantitative HCV RNA value within the last 6 months
 - Fibrosis stage must be provided

Clinical Notes:

1. Treatment-experienced is defined as a patient who has been previously treated with a peginterferon/ribavirin regimen, including regimens containing HCV protease inhibitors and who has not experienced an adequate response.
2. Acceptable methods for the measurement of fibrosis score include Fibrotest, liver biopsy, transient elastography (FibroScan®), serum biomarker panels (such as AST-to-Platelet Ratio Index or Fibrosis-4 score) either alone or in combination.
3. Compensated cirrhosis is defined as a CTP score of 5 to 6 (Class A) and decompensated cirrhosis as a CTP

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

score of 7 or above (Class B or C).

4. Re-treatment for direct-acting antiviral failures will be considered on a case-by-case basis.

Claim Notes:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904233
 - 00904234
- Claims will be limited to a 28-day supply.

** [Criteria Code 34] has been added for use effective December 1, 2019 to allow payment of a patient's initial 28 day supply only. Criteria code 34 should be provided by the prescribing physician only.

SOFOSBUVIR/VELPATASVIR/VOXILAPREVIR (Vosevi 400mg/100mg/100mg Tablet)

- For treatment-experienced adult patients with chronic hepatitis C virus (HCV) who meet the following criteria:

Approval Period

Genotypes 1, 2, 3, 4, 5, 6 or mixed genotypes

- | | |
|------------------------------|------------|
| ▪ With compensated cirrhosis | ▪ 12 weeks |
| ▪ With no cirrhosis | |

- **Patients must also meet all of the following criteria:**
 - Must be prescribed by a hepatologist, gastroenterologist, or infectious disease specialist (or other physician experienced in treating a patient with hepatitis C infection)
 - Lab-confirmed hepatitis C genotype 1, 2, 3, 4, 5, 6 or mixed genotypes
 - Quantitative HCV RNA value within the last 6 months
 - Fibrosis stage must be provided

Clinical Notes:

1. Treatment experienced is defined as a patient who has been previously treated with an NS5A inhibitor for genotype 1, 2, 3, 4, 5 or 6 or sofosbuvir without an NS5A inhibitor for genotype 1, 2, 3 or 4 and who has not experienced an adequate response.
2. Compensated cirrhosis is defined as a CTP score of 5 to 6 (Class A).
3. Re-treatment for sofosbuvir-velpatasvir-voxilaprevir treatment failures will be considered on a case-by-case basis.

Claim Notes:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904312
 - 00904313
- Claims will be limited to a 28-day supply.

** [Criteria Code 34] has been added for use effective December 1, 2019 to allow payment of a patient's initial 28 day supply only. Criteria code 34 should be provided by the prescribing physician only.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

SOMATROPIN (*Humatrope, Genotropin GoQuick and MiniQuick, Norditropin FlexPro 5mg/1.5mL, 10mg/1.5mL and 15mg/1.5mL Prefilled Pen, Nutropin, Nutropin AQ, Nutropin AQ NuSpin, Saizen Injection and Cartridge*)

- For treatment of growth hormone deficiency in patients with Turner Syndrome, upon the request of an endocrinologist or prescriber with a specialty in endocrinology

Note:

- The larger 8.8mg/vial format can be approved when suitable for dosing requirements, if it does not result in drug wastage.

***SORAFENIB** (*Nexavar 200mg Tablet*)

HEPATOCELLULAR CARCINOMA (HCC)

- For the treatment of adult patients with a diagnosis of hepatocellular carcinoma (HCC) as either first line-treatment, or second-line treatment following atezolizumab in combination with bevacizumab, who meet all the following criteria:
 - Child-Pugh Class A liver dysfunction (mild hepatic impairment)
 - ECOG performance status of 0 or 1
 - Who have either progression of disease, or who are not candidates for curative intent treatments (transplantation, hepatic resection), or other well established palliative interventions (ablation, transcatheter arterial chemo-embolization (TACE), internal radiation)

Clinical Note:

- Patients who are unable to tolerate sorafenib may be switched to lenvatinib if there is no disease progression and provided all other funding criteria are met.
- Patients with disease progression on sorafenib are not eligible for reimbursement of lenvatinib.

METASTATIC RENAL CELL CARCINOMA

- For the treatment of patients with advanced or metastatic renal cell carcinoma when used as a second-line therapy following disease progression on cytokine therapy.

Clinical Notes:

- Patients must have a good performance status.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.

STIRIPENTOL (*Diacomit 250mg and 500mg Capsule, 250mg and 500mg Powder for Suspension*)

- For use in combination with clobazam and valproate as adjunctive therapy of refractory generalized tonic-clonic seizures in patients with severe myoclonic epilepsy in infancy (Dravet syndrome), whose seizures are not adequately controlled with clobazam and valproate alone.
- The patient must be under the care of a neurologist or a pediatrician.

SUCROFERRIC OXYHYDROXIDE (*Velphoro 500mg Tablet*)

- For the treatment of hyperphosphatemia (>1.8 mmol/L) in patients with end-stage renal disease (eGFR < 15 mL/min) who have:
 - Inadequate control of phosphate levels on a calcium based phosphate binder, OR
 - Hypercalcemia (corrected for albumin), OR
 - Calciphylaxis (calcific arteriopathy)

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claim Notes:

- Must be prescribed by a nephrologist or other prescriber within the Provincial Dialysis Program.
- Initial Approval: 6 months.
- Renewal Approval: 1 year. Confirmation of improvement of phosphate levels is required (lab values must be provided).

SUMATRIPTAN (*Imitrex 50mg, 100mg Tablet and generic brands*)

- See [Selective 5HT₁ - Receptor Agonists](#)

SUMATRIPTAN (*Imitrex 6mg/Syringe Injection and generic brands*)

- See [Selective 5HT₁ - Receptor Agonists](#)

***SUNITINIB** (*Sutent 12.5mg, 25mg, 50mg Capsule and generic brands*)**METASTATIC RENAL CELL CARCINOMA**

- For patients with advanced or metastatic renal cell carcinoma as either first-line therapy, or second-line therapy after failure of first-line immunotherapy.

Clinical Notes:

- Patients must have a good performance status.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.
- Sunitinib may not be used after another tyrosine kinase inhibitor (i.e., sorafenib, or pazopanib) as sequential therapy.
- In the event of significant toxicity, a switch to another tyrosine kinase inhibitor (i.e., sorafenib or pazopanib) may be allowed.
- Both clear cell and non-clear cell histology are eligible for treatment.

ADVANCED GASTROINTESTINAL STROMAL TUMOR (GIST)

- As a single agent for the treatment of advanced gastrointestinal stromal tumor (GIST) patients after failure of imatinib due to intolerance or resistance

ADVANCED OR METASTATIC PANCREATIC NEUROENDOCRINE TUMORS (PNET)

- For the treatment of patients with progressive, unresectable, well or moderately differentiated, locally advanced or metastatic pancreatic neuroendocrine tumors (pNET) with good performance status (ECOG 0-2), until disease progression

TACROLIMUS (*Protopic 0.03%, 0.1% Ointment*)

- For children greater than 2 years of age with refractory atopic dermatitis. Coverage will be renewed yearly
- For the intermittent use for moderate to severe atopic dermatitis in adults who have:
 - failed or are intolerant to a site appropriate strength of corticosteroid therapy (i.e., low potency on face versus intermediate to high potency for trunk and extremities)

TAFAMIDIS (*Vyndamax 61mg Capsule*)**TAFAMIDIS MEGLUMINE** (*Vyndaqel 20mg Capsule*)

- For the treatment of cardiomyopathy in adult patients with documented hereditary or wild-type transthyretin-mediated amyloidosis (ATTR) who meet all of the following criteria:
 - New York Heart Association (NYHA) class I to III heart failure

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- At least one prior hospitalization for heart failure or clinical evidence of heart failure that required treatment with a diuretic
- Has not previously undergone a heart or liver transplant
- Does not have an implanted cardiac mechanical assist device (CMAD)

Discontinuation Criteria:

- The patient has:
 - NYHA class IV heart failure, or
 - received an implanted CMAD, or
 - received a heart or liver transplant.

Clinical Notes:

1. Wild-type ATTR-cardiomyopathy (CM) consists of all of the following:
 - a. absence of a variant transthyretin (TTR) genotype
 - b. TTR precursor protein identification by immunohistochemistry, scintigraphy, or mass spectrometer
 - c. evidence of cardiac involvement by echocardiography with end-diastolic interventricular septal wall thickness greater than 12 mm
 - d. presence of amyloid deposits in biopsy tissue (fat aspirate, salivary gland, median nerve connection tissue sheath, or cardiac tissue) or positive findings on technetium-99m pyrophosphate (Tc-99m-PYP) scintigraphy with single-photon emission computed tomography (SPECT) scanning
2. Hereditary ATTR-CM consists of all of the following:
 - a. presence of a variant TTR genotype associated with CM and presenting with a CM phenotype
 - b. evidence of cardiac involvement by echocardiography with end-diastolic interventricular septal wall thickness greater than 12 mm
 - c. presence of amyloid deposits in biopsy tissue (fat aspirate, salivary gland, median nerve connective tissue sheath, or cardiac tissue) or positive findings on Tc-99m-PYP scintigraphy with SPECT scanning

Claim Notes:

- The patient must be under the care of a physician with experience in the diagnosis and treatment of ATTR-CM.
- Combination therapy with other interfering ribonucleic acid drugs or transthyretin stabilizers used to treat ATTR will not be reimbursed.
- Claims will be limited to a 30-day supply.
- Initial approval period: 9 months.
- Renewal approval period: 1 year.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - Vyndamax 00904778
 - Vyndaqel 00904637

TALIGLUCERASE ALFA (*Ellyso 200U Vial*)

- For the treatment of patients with symptomatic Gaucher disease type 1 (GD1) for whom treatment with velaglucerase alfa is not tolerated or contraindicated.

Clinical Notes:

- Velaglucerase alfa is the preferred reimbursed enzyme replacement therapy for GD1.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Requests for patients currently using taliglucerase alfa who do not have a contraindication or intolerance to velaglucerase alfa will be considered for coverage of velaglucerase alfa only.
- Requests for coverage must meet the criteria for diagnosis of GD1, indication for therapy and expected response to enzyme replacement therapy outlined below:

Initial Coverage

Diagnosis:

- The diagnosis of GD1 must have been established by the demonstration of specific deficiency of glucocerebrosidase (GCase) in tissue or cultured skin fibroblasts, or by demonstration of the presence, in tissue or peripheral blood leukocytes, of mutations in the GCase gene known to result in severe enzyme deficiency.
- Other potentially confounding diagnoses, such as Hodgkin disease or other storage disorders, must have been ruled out. The symptoms experienced by the patient should be shown to be attributable to GD1 and not another condition that might mimic it.
- The patient should not have any GD1-related or other medical condition that might reasonably be expected to compromise their response to treatment. In some patients with GD1, secondary pathologic changes, such as avascular necrosis of bone, may already have occurred that would not be expected to respond to enzyme replacement. In such patients, reversal of the pathology is unlikely.

Disease Severity

Evidence of disease severity must be provided, and include at least one of the following:

- **Hematological complications**
 - Hemoglobin <85% of lower limit of age- and sex-appropriate normal after other causes of anemia, such as iron deficiency, have been treated or ruled out.
 - Platelet count <50 x 10⁹/L on two separate occasions at least one month apart. Higher cut offs may be considered in the event the patient is symptomatic with bleeding or bruising.
 - At least two episodes of severely symptomatic splenic infarcts confirmed by CT or other imaging of the abdomen.
- **Skeletal complications**
 - A single acute bone crisis severe enough to require hospitalization or marked incapacitation.
 - Radiographic or MRI evidence of incipient destruction of any major joint (e.g., hips and shoulders) or significant worsening of bony pathology (e.g. marrow infiltration, avascular necrosis, and infarcts).
 - Spontaneous fractures with evidence from imaging studies that recurrence is likely.
 - Chronic bone pain causing significant loss of time from work or school and not controlled by administration of non-narcotic analgesics or anti-inflammatory drugs.
 - Note: Patients who are scheduled for major joint replacement surgery, made necessary by skeletal complications of GD1, should be treated with enzyme therapy at a dosage of at least 30 units/kg every 2 weeks for at least 6 months before the joint replacement surgery and the dose continued until rehabilitation from the surgery is complete.
- **Gastrointestinal complications**
 - Evidence of significant liver dysfunction attributable to GD1, such as portal hypertension or impaired hepatic synthetic function. Elevation of transaminase levels with no evidence of portal hypertension or impairment in synthetic function is not an indication for ERT.
 - Significant discomfort due to enlargement of the spleen or liver.
- **Pulmonary complications**
 - Evidence of clinically significant and/or progressive pulmonary disease due to GD1.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- **Systemic complications**

- Growth failure in children: significant decrease in percentile linear growth over a 3 - 6 month period.

Exclusion Criteria:

- Due to the absence of data demonstrating therapy of asymptomatic patients alters long term outcomes, asymptomatic patients will not be considered for coverage.
- Data does not suggest that ERT is effective in improving central nervous system involvement in patients with Type 2 and 3 disease. Therefore, patients exhibiting primary neurological disease due to GD1 will not be considered for coverage. Treatment for patients at risk of neuronopathic disease should be guided by the non-neurological manifestations of their disease as outlined above and ERT should not be initiated in asymptomatic patients who have a genotype that increases their risk of neuronopathic involvement.

Continued Coverage:

- Patients' disease severity must be re-assessed annually.
- A patient may receive approval for further coverage for treatment where there is demonstrated clinical improvement based on the expected response outlined below:

Indication for therapy	Expected Response
Hemoglobin < 85% of lower limit of age and sex-appropriate normal	Increase hemoglobin levels to > 110 for women and children and > 120 for men
Platelet count < 50 x 10 ⁹ /L on two separate occasions, or bleeding complications associated with thrombocytopenia irrespective of the platelet count	Increase platelet count to level sufficient to prevent spontaneous bleeding Normalization of platelet count in splenectomized patients In patients with intact spleen, an increase of at least 1.5X baseline value
Two episodes of severely symptomatic splenic infarcts	Reduction of spleen volume by 50% Prevention of further splenic infarcts
Acute bone crises	Prevent bone crises
Radiographic or MRI evidence of incipient destruction of any major joint	Improvement in imaging parameters (either MRI, QCSI ¹ , or BMD)
Spontaneous fractures	Prevention of further fractures
Chronic bone pain	Reduce bone pain
Major joint replacement surgery	Optimize surgical outcome
Significant hepatic dysfunction	Improvement in hepatic function
Symptomatic hepatosplenomegaly	Reduction of spleen volume by 50% Reduction in liver volume by 30%
Progressive pulmonary disease due to GD1	Improvement in pulmonary hypertension ² Improvement in oxygenation Reversal of hepatopulmonary syndrome
Growth failure in children	Return to normal range of growth parameters

¹ QCSI- quantitative chemical shift imaging

² May require adjuvant treatment for pulmonary hypertension

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Discontinuation of Coverage:

- Renewals will NOT be approved if:
 - The patient or the patient's specialist fails to comply adequately with treatment or measures taken to evaluate the effectiveness of the therapy (e.g. monitoring for expected response).
 - Therapy fails to relieve the symptoms of disease that originally resulted in the patient being approved for treatment.

Claim Notes:

- Approvals will be for a maximum of 60 units/kg every 2 weeks.
- Initial Approval: 6 months.
- Renewal Approval: 1 year.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904383
 - 00904385

TAZAROTENE (*Arazlo 0.045% Lotion*)

- Regular benefit for beneficiaries 30 years and under
- For treatment of acne vulgaris in beneficiaries over the age of 30

TEDUGLUTIDE (*Revestive 5mg Powder for Injection*)

ADULT PATIENTS WITH SHORT BOWEL SYNDROME (SBS)

- **For the treatment of adult patients with Short Bowel Syndrome (SBS) who have all of the following:**
 - SBS as a result of major intestinal resection (e.g., volvulus, vascular disease, cancer, Crohn's disease, injury).
 - dependency on parenteral nutrition (PN) for a least 12 months.
 - prior to initiating teduglutide, PN required at least three times weekly to meet caloric, fluid or electrolyte needs, due to ongoing malabsorption and stable PN frequency and volume for at least one month.

Renewal Criteria:

- Has maintained at least a 20% reduction in PN volume from baseline at 12 months.

Clinical Note:

- PN is defined as the parenteral delivery of lipids, protein and/or carbohydrates to address caloric needs, and intravenous fluids which addresses fluid and electrolyte needs of patients.

Claim Notes:

- Must be prescribed by a specialist with experience in SBS.
- Approval period: 1 year.

PEDIATRIC PATIENTS 1 YEAR OF AGE AND OLDER WITH SHORT BOWEL SYNDROME (SBS)

- **For the treatment of pediatric patients 1 year of age and older with Short Bowel Syndrome (SBS) who have all of the following:**
 - Prior to initiating teduglutide, parenteral support (PS) requirements must be stable or there must have been no improvement in enteral feeding for at least three months.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- PS must provide more than 30% of caloric and/or fluid/electrolyte needs.
- The cumulative lifetime duration of PS must be at least 12 months.

Renewal Criteria:

- Has maintained at least a 20% reduction in parenteral support volume from baseline.

Claim Notes:

- Must be prescribed by a pediatric gastroenterologist or other prescriber currently working within a specialized multi-disciplinary intestinal rehabilitation program with expertise in the diagnosis and management of SBS.
- Initial approval period: 6 months.
- Renewal approval period: 6 months.

Clinical Note:

- PS is defined as the parenteral delivery of lipids, protein and/or carbohydrates to address caloric needs, and intravenous fluids which addresses fluid and electrolyte needs of patients.

***TEMOZOLOMIDE** (*Temodal 5mg, 20mg, 100mg, 140mg, 250mg Capsule and generic brands*)

- For the treatment of patients with high grade gliomas as monotherapy or in combination with other therapies such as radiation.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be continued until there is no longer a clinical benefit or unacceptable toxicity.

TERIFLUNOMIDE (*Aubagio 14mg Tablet and generic brands*)

- For the treatment of patients with relapsing remitting multiple sclerosis (RRMS) who meet all of the following criteria:
 - requested and followed by a neurologist experienced in the management of RRMS; and
 - recent expanded disability status scale (EDSS) score of 5.5 or less (i.e. patients must be able to ambulate at least 100 metres without assistance).
- **Exclusions:**
 - not funded in combination with other disease modifying therapies;
 - not funded in patients with an EDSS > 5.5;
 - not funded in patients < 18 years of age.
- **Renewals:**
 - EDSS score < 5.5 (i.e. patients must be able to ambulate at least 100 metres without assistance). Date and details of the most recent neurological examination and EDSS score must be provided (exam must have occurred within the last 90 days); and
 - Patients must be stable or have experienced no more than 1 disabling attack/relapse in the past year.

***TESTOSTERONE, TOPICAL** (*Androgel Gel Packet and generic brands, and Testim Gel*)

- For the treatment of congenital and acquired primary or secondary hypogonadism in males with a specific diagnosis of:
 - Primary: cryptorchidism, Klinefelter's, orchidectomy, and other established causes; OR
 - Secondary: pituitary-hypothalamic injury due to tumors, trauma, radiation; AND

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- For those with one of the above diagnoses, the deficiency must be clearly demonstrated by clinical features and confirmed by two separate biochemical tests.

Notes:

- Maximum dose approved is 5g per day.
- This will be adjudicated by limiting the quantity payable each quarter (e.g. Jan-Mar) to:
 - 300g of Androgel 2.5g gel (packet);
 - 600g of Androgel 5g gel (packet); or
 - 600g of Testim Gel.

Please be reminded that topical gels are to be billed per gram (not per packet).

TESTOSTERONE, TOPICAL (*Androgel Gel Packet and generic brands, and Testim Gel*)

- For use in gender affirming hormone therapy.

Claim Note:

- Maximum dose approved is 5g gel per day.

TEZPELUMAB (*Tezspire 210mg/1.91mL Prefilled Pen and Syringe*)

- For the treatment of severe asthma in patients 12 years and older who are inadequately controlled with high-dose inhaled corticosteroids (ICS), and one or more additional asthma controller(s) (e.g., long-acting beta-agonist), and have experienced 2 or more clinically significant asthma exacerbations in the past 12 months.

Initial Discontinuation Criteria:

- Baseline asthma control questionnaire score has not improved at 12 months since initiation of treatment, or
- No decrease in the daily maintenance OCS dose in the first 12 months of treatment, or
- The number of clinically significant asthma exacerbations has increased within the previous 12 months.

Subsequent Discontinuation Criteria:

- Asthma control questionnaire score achieved after the first 12 months of therapy has not been maintained subsequently, or
- The reduction in the daily maintenance dose of OCS achieved after the first 12 months of treatment is not maintained or improved subsequently, or
- The number of clinically significant asthma exacerbations has increased within the previous 12 months.

Clinical Notes:

- A baseline assessment of asthma symptom control using a validated asthma control questionnaire must be provided.
- A baseline and annual number of clinically significant asthma exacerbations must be provided.
- High dose ICS is defined as ≥ 500 mcg of fluticasone propionate or equivalent daily dose.
- A significant clinical exacerbation is defined as worsening of asthma such that the treating physician elected to administer systemic glucocorticoids for at least 3 days or the patient visited an emergency department or was hospitalized.

Claim Notes:

- Must be prescribed by a respirologist, clinical immunologist, allergist or internist experienced in the treatment of

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

severe asthma.

- Combined use of tezepelumab with other biologics used to treat asthma will not be reimbursed.
- Approvals will be for a maximum of 210mg subcutaneous injection every 4 weeks.
- Approval period: 1 year.

***THYROTROPIN** (*Thyrogen 0.9mg/mL Injection*)

- To monitor for recurrence and metastatic disease, in patients who have documented evidence of thyroid cancer and who have undergone appropriate surgical and/or medical management. This includes:
 - primary use in patients with inability to raise an endogenous TSH level (≥ 25 mu/L) with thyroid hormone withdrawal;
 - primary use in cases of documented morbidity in patients for whom severe hypothyroidism could be life-threatening;
 - secondary use in patients with previous thyroid hormone withdrawal resulting in a documented life-threatening event.
- As a single agent for the preparation of radioiodine remnant ablation in patients with papillary or follicular thyroid cancer who have undergone thyroidectomy as treatment for thyroid cancer
 - thyrotropin is a reasonable alternate to thyroid hormone withdrawal in patients who are unable to tolerate the prolonged hypothyroid state or who cannot achieve satisfactory elevation of endogenous TSH;
 - thyrotropin may be used in new patients or patients with previously incomplete remnant ablation or who have a recurrence of thyroid cancer and require therapeutic remnant ablation.

TICLOPIDINE (*Ticlid 250mg Tablet and generic brands*)

- For the secondary prevention of ischemic stroke or transient ischemic attack (TIA) in patients with a documented severe allergy to ASA or who experience a recurrent thrombotic event (stroke, symptoms of TIA) while taking ASA. **[Criteria Code 01]**
- For the prevention of thrombosis in patients post intracoronary stent implantation for a period of up to 30 days following insertion. **[Criteria Code 02]**
- Other requests on a case by case basis.

TICAGRELOR (*Brilinta 90mg Tablet and generic brands*)

- To be taken in combination with ASA 75 mg -150mg daily¹ for patients with acute coronary syndrome (i.e. ST elevation myocardial infarction (STEMI), non-ST elevation myocardial infarction (NSTEMI), or unstable angina (UA), as follows:

STEMI^{2,3}

- STEMI patients undergoing primary percutaneous coronary intervention (PCI).

NSTEMI or UA^{2,3}

- presence of high risk features irrespective of intent to perform revascularization:
 - high GRACE risk score (>140);
 - high TIMI risk score (5-7);
 - second ACS within 12 months;
 - complex or extensive coronary artery disease e.g. diffuse three vessel disease;
 - definite documented cerebrovascular or peripheral vascular disease;
 - previous CABG;

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

OR

- undergoing PCI + high risk angiographic anatomy⁴.
- Coverage duration: 12 months.

Note:

- **[Criteria Code 30]** may be used for the initial 30 day coverage period, however a written request submitted to the Pharmacare office is required to allow coverage for the remaining duration of treatment.
1. Co-administration of ticagrelor with high maintenance dose ASA (>150 mg daily) is not recommended.
 2. In the PLATO study more patients on ticagrelor experienced non CABG related major bleeding than patients on clopidogrel, however, there was no difference between the rate of overall major bleeding, between patients treated with ticagrelor and those treated with clopidogrel. As with all other antiplatelet treatments the benefit/risk ratio of antithrombotic effect vs. bleeding complications should be evaluated.
 3. Ticagrelor is contraindicated in patients with active pathological bleeding, in those with a history of intracranial hemorrhage and moderate to severe hepatic impairment.
 4. High risk angiographic anatomy is defined as any of the following: left main stenting, high risk bifurcation stenting (i.e., two-stent techniques), long stents ≥ 38 mm or overlapping stents, small stents ≤ 2.5 mm in patients with diabetes.

***TIGECYCLINE** (*Tygacil 50mg Vial*)

- For the treatment of patients with multi-drug resistant infections when alternative agents are not an option.

Claim Note:

- Must be prescribed by, or in consultation with, an infectious disease specialist or medical microbiologist.

TILDRAKIZUMAB (*Ilumya 100 mg/mL Prefilled Syringe*)

- For patients with severe, debilitating chronic plaque psoriasis who meet all of the following:
 - Body surface area (BSA) involvement of >10% and/or significant involvement of the face, hands, feet or genitals;
 - Failure to, contraindication to or intolerant of methotrexate and cyclosporine;
 - Failure to, intolerant of or unable to access phototherapy;
 - Written request of a dermatologist or prescriber with a specialty in dermatology.
- Continued coverage is dependent on evidence of improvement, specifically:
 - A >75% reduction in the Psoriasis Area and Severity Index (PASI) score; or
 - A >50% reduction in PASI with a > 5 point improvement in DLQI (Dermatology Life Quality Index); or
 - Significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals.

Clinical Note:

- Treatment should be discontinued if a response has not been demonstrated after 16 weeks.

Claim Notes:

- Concurrent use of biologics not approved.
- Approvals will be for 100 mg by subcutaneous injection at week 0, week 4, and every 12 weeks thereafter.
- Initial approval period: 16 weeks
- Renewal approval period: 1 year

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

TIOTROPIUM BROMIDE (*Spiriva 18mcg Cap for Inhalation*)

TIOTROPIUM BROMIDE MONOHYDRATE (*Spiriva Respimat 2.5µg/actuation Inhaled Solution*)

- See [Long-Acting Muscarinic Antagonists](#)

TIOTROPIUM BROMIDE MONOHYDRATE/OLODATEROL HYDROCHLORIDE (*Inspiroto Respimat 2.5mcg/2.5mc g Inhaled Solution*)

- See [Long-Acting Beta₂-Agonists/Long-Acting Muscarinic Antagonists](#)

***TRIFLURIDINE AND TIPIRACIL** (*Lonsurf 15mg/6.14mg and 20mg/8.19mg Tablet*)

- For the treatment of adult patients with metastatic gastric cancer or adenocarcinoma of the gastroesophageal junction who meet the following criteria:
 - Previously treated with at least two prior lines of chemotherapy including a fluoropyrimidine, a platinum, and either a taxane or irinotecan and if appropriate, with HER2-targeted therapy.
 - Patients should have a good performance status.

Clinical Notes:

- Trifluridine/tipiracil should be used in combination with best supportive care.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.
- Requests will be considered for patients who have an intolerance or contraindication to platinum-based therapy.

TOCILIZUMAB (*Actemra 80mg/4mL, 200mg/10mL, 400mg/20mL Injection, 162mg/0.9mL SC Injection and Autoinjector*)

GIANT CELL ARTERITIS (GCA) (ACTEMRA 162MG/0.9ML SC INJECTION AND AUTOINJECTOR)

- For the treatment of Giant Cell Arteritis (GCA) in adult patients who are receiving prednisone at initiation of therapy, or with relapse.

Notes:

- Patients should be under the care of a physician with the experience of diagnosis and management of GCA.
- Duration of therapy with tocilizumab should be limited to 52 weeks per treatment course.
- Discontinuation of tocilizumab should be considered at 12 weeks if there is no response to therapy.

POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS (PJIA) (ACTEMRA 80MG/4ML, 200MG/10ML, 400MG/20ML INJECTION AND 162MG/0.9ML SC INJECTION AND AUTOINJECTOR)

- For the treatment of children (age 2-17) with moderately to severely active polyarticular juvenile idiopathic arthritis (pJIA) who have had inadequate response to one or more disease-modifying antirheumatic drugs (DMARDs).

Notes:

- Must be prescribed by, or in consultation with, a rheumatologist who is familiar with the use of biologic DMARDs in children.
- Intravenous infusion: Approvals will be for 10mg/kg for patients <30kg or 8mg/kg for patients ≥ 30kg, to a maximum of 800mg, administered every four weeks.
- Subcutaneous injection: Approvals will be for a maximum of 162mg once every three weeks for patients weighing <30kg or 162mg once every two weeks for patients weighing ≥30kg.
- Initial approval period: 16 weeks
- Renewal Approval: 1 year. Confirmation of continued response is required.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

RHEUMATOID ARTHRITIS (RA) (ACTEMRA 80MG/4ML, 200MG/10ML, 400MG/20ML INJECTION AND 162MG/0.9ML SC INJECTION AND AUTOINJECTOR)

- For the treatment of severely active rheumatoid arthritis, in combination with methotrexate or other disease-modifying antirheumatic drugs (DMARDs), in adult patients who are refractory or intolerant to:
 - methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age), or use in combination with another DMARD, for a minimum of 12 weeks

AND

- methotrexate in combination with at least two other DMARDs, such as hydroxychloroquine and sulfasalazine, for a minimum of 12 weeks

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- Optimal treatment response to DMARDs may take up to 24 weeks, however coverage of a biologic therapy can be considered if no improvement is seen after 12 weeks of triple DMARD use.
- If patient factors (e.g. intolerance) prevent the use of triple DMARD therapy, these must be described and dual therapy with DMARDs must be tried.
- 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. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Combined use of more than one biologic DMARD will not be reimbursed.
- Initial Approval: 6 months.
- Renewal Approval: 1 year. Confirmation of continued response is required.
- **Maximum Dosage Approved:**
 - Tocilizumab: 4mg/kg/dose once every 4 weeks followed by an increase to 8 mg/kg/dose based on clinical response

SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS (SJIA) (ACTEMRA 80MG/4ML, 200MG/10ML, 400MG/20ML INJECTION AND 162MG/0.9ML SC INJECTION AND AUTOINJECTOR)

- For the treatment of active systemic juvenile idiopathic arthritis (sJIA), in patients 2 years of age or older, who have responded inadequately to non-steroidal anti-inflammatory drugs (NSAIDs) and systemic corticosteroids (with or without methotrexate) due to intolerance or lack of efficacy.

Notes:

- Must be prescribed by, or in consultation with, a rheumatologist, who is familiar with the use of biologic DMARDs in children.
- Intravenous infusion: Approvals will be for 12 mg/kg for patients < 30 kg or 8 mg/kg for patients ≥ 30 kg, to a maximum of 800mg, administered every two weeks.
- Subcutaneous injection: Approvals will be for a maximum of 162mg once every two weeks for patients weighing < 30 kg or 162mg once every week for patients weighing ≥ 30 kg.
- Initial approval period: 16 weeks
- Renewal Approval: 1 year. Confirmation of continued response is required.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

TOFACITINIB (*Xeljanz 5mg, 10mg Tablet and Xeljanz XR 11mg Tablet*)

RHEUMATOID ARTHRITIS (XELJANZ 5MG TABLET AND XELJANZ XR 11MG TABLET)

- For the treatment of severely active rheumatoid arthritis, in combination with methotrexate or other disease-modifying antirheumatic drugs (DMARDs), in adult patients who are refractory or intolerant to:
 - methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age) for a minimum of 12 weeks, followed by methotrexate in combination with at least two other DMARDs, such as hydroxychloroquine and sulfasalazine, for a minimum of 12 weeks;
OR
 - initial use of triple DMARD therapy with methotrexate in combination with at least two other DMARDs such as hydroxychloroquine and sulfasalazine, for a minimum of 24 weeks.

Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- Optimal treatment response may take up to 24 weeks; however coverage of tofacitinib can be considered if no improvement is seen after 12 weeks of triple DMARD use.
- If the patient is intolerant to triple DMARD therapy, then dual therapy with DMARDs (methotrexate, hydroxychloroquine, leflunomide, sulfasalazine) must be considered.
- 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. The nature of intolerance(s) must be clearly documented.
- Must be prescribed by a rheumatologist.
- Combined use with biologic DMARD will not be reimbursed.

ULCERATIVE COLITIS (XELJANZ 5MG AND XELJANZ 10MG TABLET)

- For the treatment of adult patients with moderately to severely active ulcerative colitis who have a partial Mayo score > 4 , and a rectal bleeding subscore ≥ 2 and are:
 - refractory or intolerant to conventional therapy (i.e. 5-ASA for a minimum of 4 weeks, and prednisone ≥ 40 mg daily for two weeks or IV equivalent for one week); OR
 - corticosteroid dependent (i.e. cannot be tapered from corticosteroids without disease recurrence; or have relapsed within three months of stopping corticosteroids; or require two or more courses of corticosteroid within one year.)
- Renewal requests must include information demonstrating the beneficial effects of the treatment, specifically:
 - a decrease in the partial Mayo score ≥ 2 from baseline, AND
 - a decrease in the rectal bleeding subscore ≥ 1 .

Clinical Notes:

- 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. The nature of intolerance(s) must be clearly documented.
- Patients with severe disease do not require a trial of 5-ASA

Claim Notes:

- Must be prescribed by a gastroenterologist or physician with a specialty in gastroenterology.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Combined use with one or more biologic DMARD will not be reimbursed.
- Approvals will be for a maximum dose of 10 mg twice daily (Xeljanz).
- Initial Approval: 16 weeks.
- Renewal Approval: 1 year.

TOPIRAMATE (*Topamax 15mg and 25mg Sprinkle Capsule*)

- For patients who require topiramate, cannot take the tablet form, and require sprinkle capsules for proper administration.

TRAMETINIB (*Mekinist 0.5mg and 2mg Tablet*)

- See [Dabrafenib \(Tafinlar 50mg and 75mg Capsule\) and Trametinib \(Mekinist 0.5mg and 2mg Tablet\)](#)

TRETINOIN (*Stieva-A, Retin-A, Vitamin A Acid Topical Preparations*)

- Regular benefit for beneficiaries 30 years and under
- For treatment of acne vulgaris in beneficiaries over the age of 30

TRIAMCINOLONE HEXACETONIDE (*Trispan 20mg/mL Injection*)

- For the treatment of juvenile idiopathic arthritis.

TRIENTINE HYDROCHLORIDE (*250mg Capsule*)

WILSON'S DISEASE

- For the treatment of Wilson's disease in patients who have experienced intolerance or have a contraindication to d-penicillamine.

Clinical Note:

- Intolerance is defined as demonstrating serious adverse effects to treatments. The nature of intolerance(s) must be clearly documented.

Claims Notes:

- Treatment must be initiated by clinicians experienced in the management of Wilson's disease for adult patients 18 years of age or older.
- Treatment must be initiated and renewed by clinicians experienced in the management of Wilson's disease for patients less than 18 years of age.
- Approval: 12 months

TRihePTANOIN (*Dojolvi 100% Oral Liquid*)

- For the treatment of adult and pediatric patients with an acute life-threatening long-chain fatty acid oxidation disorders (LC-FAOD) who meet the following criteria:
 - patients with a confirmed diagnosis of LC-FAOD and acute life-threatening events who require alternative therapy to conventional even-chain medium-chain triglyceride (MCT) supplementation, OR
 - patients without a confirmed diagnosis of LC-FAOD presenting with acute life-threatening events consistent with LC-FAOD who require alternative therapy to conventional even-chain MCT supplementation.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Claims Notes:

- Triheptanoin should only be prescribed by clinicians experienced in the management of LC-FAOD.
- Approval: 1 year. Confirmation of continued response required.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - 00900021

TROSPIUM (*Trosec 20mg Tablet and generic brands*)

- See [OAB Medications](#)

TRYPTOPHAN (*Tryptan 500mg Capsule and 500mg, 750mg, 1g Tablet and generic brands*)

- As an adjunct for the treatment of depression in the management of patients suffering from bipolar affective disorders

***TUCATINIB** (*Tukysa 50mg and 150mg Tablet*)**LOCALLY ADVANCED UNRESECTABLE OR METASTATIC HER2-POSITIVE BREAST CANCER**

- In combination with trastuzumab and capecitabine for the treatment of patients with locally advanced unresectable or metastatic HER2-positive breast cancer who have received prior treatment with trastuzumab, pertuzumab and a HER2-targeted antibody-drug conjugate (e.g., trastuzumab emtansine or trastuzumab deruxtecan), where at least one was given in the advanced or metastatic setting.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be discontinued upon disease progression, unacceptable toxicity, or if both trastuzumab and capecitabine are discontinued.

Claim Note:

- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PIN:
 - Tukysa 150mg Tab 00904820

UMECLIDINIUM (AS BROMIDE) (*Incruse Ellipta 62.5mcg Dry Powder for Oral Inhalation*)

- See [Long-Acting Muscarinic Antagonists](#)

UMECLIDINIUM (AS BROMIDE) AND VILANTEROL (AS TRIFENATATE) (*Anoro Ellipta 62.5mcg/25mcg Powder for Oral Inhalation*)

- See [Long-Acting Beta₂-Agonists/Long-Acting Muscarinic Antagonists](#)

UPADACITINIB (*Rinvoq 15mg and 30mg Tab*)**RHEUMATOID ARTHRITIS**

- For the treatment of severely active rheumatoid arthritis, alone or in combination with methotrexate or other disease-modifying antirheumatic drugs (DMARDs), in adult patients who are refractory or intolerant to:
 - methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age), or use in combination with another DMARD, for a minimum of 12 weeks;
- AND

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- methotrexate in combination with at least two other DMARDs, such as hydroxychloroquine and sulfasalazine, for a minimum of 12 weeks.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- Optimal treatment response to DMARDs may take up to 24 weeks, however coverage can be considered if no improvement is seen after 12 weeks of triple DMARD use.
- If patient factors (e.g. intolerance) prevent the use of triple DMARD therapy, these must be described and dual therapy with DMARDs must be tried.
- 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. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of more than one biologic DMARD or janus kinase inhibitors will not be reimbursed.
- Approvals will be for a maximum of 15 mg daily.
- Initial Approval: 6 months
- Renewal Approval Period: 1 year. Confirmation of continued response required.

PSORIATIC ARTHRITIS

- For the treatment of patients with predominantly axial psoriatic arthritis who are refractory, intolerant or have contraindications to the sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each.
- For the treatment of patients with predominantly peripheral psoriatic arthritis who are refractory, intolerant or have contraindications to:
 - The sequential use of at least two NSAIDs at maximal tolerated dose for a minimum of two weeks each; AND
 - Methotrexate (oral or parenteral) at a dose of ≥ 20 mg weekly (≥ 15 mg if patient is ≥ 65 years of age) for a minimum of 8 weeks; AND
 - Leflunomide for a minimum of 10 weeks or sulfasalazine for a minimum of 3 months.

Clinical Notes:

- For patients who do not demonstrate a clinical response to oral methotrexate, or who experience gastrointestinal intolerance, a trial of parenteral methotrexate must be considered.
- 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 to treatments. The nature of intolerance(s) must be clearly documented.

Claim Notes:

- Must be prescribed by a rheumatologist.
- Concurrent use of biologics not approved.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Approvals will be for a maximum of 15mg daily.
- Initial coverage period: 6 months.
- Renewal approval: 1 year. Confirmation of continued response required.

ATOPIC DERMATITIS

- For the treatment of moderate to severe atopic dermatitis in patients 12 years of age and older who meet all of the following criteria:
 - Patients must have had an adequate trial (with a documented refractory disease), or were intolerant (with documented intolerance), or are ineligible for each of the following therapies:
 - maximally tolerated medical topical therapies for AD combined with phototherapy (where available), and;
 - maximally tolerated medical topical therapies for AD combined with at least 1 of the 4 systemic immunomodulators (methotrexate, cyclosporine, mycophenolate mofetil, or azathioprine)
 - Baseline Physician Global Assessment score of 3 or greater and Eczema Area and Severity Index (EASI) of 7.1 or greater.

Renewal Criteria:

- Requests for renewal must provide proof of beneficial clinical effect when defined as a 75% or greater improvement from baseline in the EASI score (EASI-75) 6 months after treatment initiation.
- Proof of maintenance of EASI-75 response from baseline must be provided for subsequent authorizations.

Clinical Note:

- Not to be used in combination with phototherapy, any immunomodulatory agents (including biologics or other janus kinase inhibitor treatment) for moderate to severe AD.

Claim Notes:

- The patient must be under the care of a dermatologist, allergist, clinical immunologist, or pediatrician who has expertise in the management of moderate to severe AD.
- Approvals will be for a maximum of 30mg once daily.
- Initial approval period: 6 months
- Renewal approval period: 1 year

USTEKINUMAB (Stelara 45mg/0.5mL and 90mg/1.0mL Injection)

- For patients with severe, debilitating chronic plaque psoriasis (PsO) who meet all of the following criteria:
 - Body Surface Area (BSA) involvement of >10% and/or significant involvement of the face, hands, feet or genitals
 - failure to respond to, contraindication to or intolerant of methotrexate and cyclosporine
 - failure to respond to, intolerant of or unable to access phototherapy
- Written request of a dermatologist or prescriber with a specialty in dermatology
- Continued coverage is dependent on evidence of improvement, specifically:
 - ≥ 75% reduction in the Psoriasis Area and Severity Index (PASI) score, *or*
 - ≥ 50% reduction in PASI with a ≥ 5 point improvement in DLQI (Dermatology Life Quality Index), *or*
 - significant reduction in BSA involved, with consideration of important regions such as the face, hands, feet or genitals

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Concurrent use of biologics not approved.
- Initial approval for a maximum of 16 weeks.
- Dosage restricted to 45mg at 0, 4 and 16 weeks, response must be assessed prior to fourth dose.
- Maintenance dosing every 12 weeks.

VALGANCICLOVIR (*Valcyte 50mg/mL Powder for Oral Solution and generic brands*)

- For the treatment of cytomegalovirus (CMV) retinitis in HIV-positive patients, upon the request of an infectious disease specialist or prescriber with a specialty in infectious disease
- For the prevention of CMV disease post solid organ transplantation in patients at high-risk (D+ / R-) (i.e., donor positive/recipient negative). Coverage will be for a maximum of 90 days
- For the treatment of patients with CMV infection who have received a solid organ transplant.

Note:

- Requests for oral suspension will be considered for patients when oral tablets are not an option.

***VANDETANIB** (*Caprelsa 100mg and 300mg Tablet*)

- For the treatment of symptomatic and/or progressive medullary thyroid cancer (MTC) in patients with unresectable locally advanced or metastatic disease. Treatment should be for patients with a good performance status and should continue until disease progression or unacceptable toxicity.

VARENICLINE (*Champix 0.5mg and 1mg Tablet, 0.5mg/1mg combopack and generic brands*)

- See [Smoking Cessation Therapies](#)

VEDOLIZUMAB (*Entyvio 300mg Vial, 108mg/0.68mL Prefilled Pen and Prefilled Syringe*)

CROHN'S DISEASE

- For patients with moderate to severely active Crohn's disease and are:
 - refractory or have contraindications to an adequate course of 5-aminosalicylic acid and corticosteroids and other immunosuppressive therapy.

Clinical Note:

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

Claim Notes:

- Must be prescribed by a gastroenterologist or physician with a specialty in gastroenterology.
- Combined use of more than one biologic DMARD will not be reimbursed.
- Intravenous infusion: Initial reimbursement is restricted to induction doses of 300mg at Weeks 0, 2 and 6. Clinical response to be assessed prior to the administration of the fourth dose.
- Subcutaneous injection: Initial reimbursement is for at least two doses of intravenous infusions of vedolizumab. Clinical response to be assessed prior to the administration of the first subcutaneous dose. Subsequent reimbursement for maintenance dosing is 108mg subcutaneously every 2 weeks.
- Initial Approval: 16 weeks
- Renewal Approval: 1 year

ULCERATIVE COLITIS

- For the treatment of adult patients with moderately to severely active ulcerative colitis who have a partial Mayo

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

score > 4, and a rectal bleeding subscore ≥ 2 and are:

- refractory or intolerant to conventional therapy (i.e. 5-ASA for a minimum of 4 weeks, and prednisone ≥ 40 mg daily for two weeks or IV equivalent for one week); or
 - corticosteroid dependent (i.e. cannot be tapered from corticosteroids without disease recurrence; or have relapsed within three months of stopping corticosteroids; or require two or more courses of corticosteroids within one year.)
- Renewal requests must include information demonstrating the beneficial effects of the treatment, specifically:
 - a decrease in the partial Mayo score ≥ 2 from baseline, and
 - a decrease in the rectal bleeding subscore ≥ 1 .

Clinical Notes:

- 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. The nature of intolerance(s) must be clearly documented.
- Patients with severe disease do not require a trial of 5-ASA.

Claim Notes:

- Must be prescribed by a gastroenterologist or physician with a specialty in gastroenterology.
- Combined use of more than one biologic DMARD will not be reimbursed.
- Intravenous infusion: Initial reimbursement is restricted to induction doses of 300mg at Weeks 0, 2 and 6. Clinical response to be assessed prior to the administration of the fourth dose.
- Subcutaneous injection: Initial reimbursement is for at least two doses of intravenous infusions of vedolizumab. Clinical response to be assessed prior to the administration of the first subcutaneous dose. Subsequent reimbursement for maintenance dosing is 108mg subcutaneously every 2 weeks.
- Initial Approval: 16 weeks
- Renewal Approval: 1 year

VELAGLUCERASE ALFA (VPRIV 400U Vial)

- For the treatment of patients with symptomatic Gaucher disease type 1 (GD1) for whom treatment with velaglucerase alfa is tolerated or not contraindicated.

Clinical Notes:

- Velaglucerase alfa is the preferred reimbursed enzyme replacement therapy (i.e. first tier) for all new and existing GD1.
- Requests for patients currently using taliglucerase alfa who do not have a contraindication or intolerance to velaglucerase alfa will be switched to velaglucerase alfa only.
- Requests for coverage must meet the criteria for diagnosis of GD1, indication for therapy and expected response to enzyme replacement therapy outlined below:

Initial Coverage

Diagnosis

- The diagnosis of GD1 must have been established by the demonstration of specific deficiency of glucocerebrosidase (GCase) in tissue or cultured skin fibroblasts, or by demonstration of the presence, in tissue or peripheral blood leukocytes, of mutations in the GCase gene known to result in severe enzyme deficiency.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Other potentially confounding diagnoses, such as Hodgkin disease or other storage disorders, must have been ruled out. The symptoms experienced by the patient should be shown to be attributable to GD1 and not another condition that might mimic it.
- The patient should not have any GD1-related or other medical condition that might reasonably be expected to compromise their response to treatment. In some patients with GD1, secondary pathologic changes, such as avascular necrosis of bone, may already have occurred that would not be expected to respond to enzyme replacement. In such patients, reversal of the pathology is unlikely.

Disease Severity

Evidence of disease severity must be provided, and include at least one of the following:

- **Hematological complications**
 - Hemoglobin <85% of lower limit of age- and sex-appropriate normal after other causes of anemia, such as iron deficiency, have been treated or ruled out.
 - Platelet count <50 x 10⁹/L on two separate occasions at least one month apart. Higher cut offs may be considered in the event the patient is symptomatic with bleeding or bruising.
 - At least two episodes of severely symptomatic splenic infarcts confirmed by CT or other imaging of the abdomen.
- **Skeletal complications**
 - A single acute bone crisis severe enough to require hospitalization or marked incapacitation.
 - Radiographic or MRI evidence of incipient destruction of any major joint (e.g., hips and shoulders) or significant worsening of bony pathology (e.g. marrow infiltration, avascular necrosis, and infarcts).
 - Spontaneous fractures with evidence from imaging studies that recurrence is likely.
 - Chronic bone pain causing significant loss of time from work or school and not controlled by administration of non-narcotic analgesics or anti-inflammatory drugs.
 - Note: Patients who are scheduled for major joint replacement surgery, made necessary by skeletal complications of GD1, should be treated with enzyme therapy at a dosage of at least 30 units/kg every 2 weeks for at least 6 months before the joint replacement surgery and the dose continued until rehabilitation from the surgery is complete.
- **Gastrointestinal complications**
 - Evidence of significant liver dysfunction attributable to GD1, such as portal hypertension or impaired hepatic synthetic function. Elevation of transaminase levels with no evidence of portal hypertension or impairment in synthetic function is not an indication for ERT.
 - Significant discomfort due to enlargement of the spleen or liver.
- **Pulmonary complications**
 - Evidence of clinically significant and/or progressive pulmonary disease due to GD1.
- **Systemic complications**
 - Growth failure in children: significant decrease in percentile linear growth over a 3 - 6 month period.

Exclusion Criteria:

- Due to the absence of data demonstrating therapy of asymptomatic patients alters long term outcomes, asymptomatic patients will not be considered for coverage.
- Data does not suggest that ERT is effective in improving central nervous system involvement in patients with Type 2 and 3 disease. Therefore, patients exhibiting primary neurological disease due to GD1 will not be considered for coverage. Treatment for patients at risk of neuronopathic disease should be guided by the non-neurological manifestations of their disease as outlined above and ERT should not be initiated in asymptomatic patients who have a genotype that increases their risk of neuronopathic involvement.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Continued Coverage:

- Patients' disease severity must be re-assessed annually.
- A patient may receive approval for further coverage for treatment where there is demonstrated clinical improvement based on the expected response outlined below:

Indication for therapy	Expected Response
Hemoglobin < 85% of lower limit of age and sex-appropriate normal	Increase hemoglobin levels to > 110 for women and children and > 120 for men
Platelet count < 50 x 10 ⁹ /L on two separate occasions, or bleeding complications associated with thrombocytopenia irrespective of the platelet count	Increase platelet count to level sufficient to prevent spontaneous bleeding Normalization of platelet count in splenectomized patients In patients with intact spleen, an increase of at least 1.5X baseline value
Two episodes of severely symptomatic splenic infarcts	Reduction of spleen volume by 50% Prevention of further splenic infarcts
Acute bone crises	Prevent bone crises
Radiographic or MRI evidence of incipient destruction of any major joint	Improvement in imaging parameters (either MRI, QCSI ¹ , or BMD)
Spontaneous fractures	Prevention of further fractures
Chronic bone pain	Reduce bone pain
Major joint replacement surgery	Optimize surgical outcome
Significant hepatic dysfunction	Improvement in hepatic function
Symptomatic hepatosplenomegaly	Reduction of spleen volume by 50% Reduction in liver volume by 30%
Progressive pulmonary disease due to GD1	Improvement in pulmonary hypertension ² Improvement in oxygenation Reversal of hepatopulmonary syndrome
Growth failure in children	Return to normal range of growth parameters

1. QCSI- quantitative chemical shift imaging

2. May require adjuvant treatment for pulmonary hypertension

Discontinuation of Coverage:

- Renewals will NOT be approved if:
 - The patient or the patient's specialist fails to comply adequately with treatment or measures taken to evaluate the effectiveness of the therapy (e.g. monitoring for expected response).
 - Therapy fails to relieve the symptoms of disease that originally resulted in the patient being approved for treatment.

Claim Notes:

- Approvals will be for a maximum of 60 units/kg every 2 weeks.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

- Initial Approval: 6 months.
- Renewal Approval: 1 year.
- Claims that exceed the maximum claim amount of \$9,999.99 must be divided and submitted as separate transactions using the DIN first and then the following PINs:
 - 00904378
 - 00904379
 - 00904380

***VEMURAFENIB** (*Zelboraf 240mg Tablet*)

- As a first line, single agent for the treatment of BRAF V600 mutation positive unresectable or metastatic melanoma in patients with an ECOG performance status (PS) of ≤ 1
- For BRAF V600 mutation positive patients who have progressed after first line treatment prior to vemurafenib availability, funding of vemurafenib as a second line agent may be considered
- For the treatment of patients with BRAF V600 mutation-positive unresectable or metastatic melanoma when used alone or in combination with cobimetinib.

Renewal Criteria:

- Written confirmation that the patient has responded to treatment and there is no evidence of disease progression.

Clinical Notes:

- Patients must have a good performance status.
- If brain metastases are present, patients should be asymptomatic or have stable symptoms.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.

Claim Note:

- Vemurafenib will not be reimbursed in patients who have progressed on BRAF and/or MEK inhibitor therapy.

***VENETOCLAX** (*Venclexta 10mg, 50mg, 100mg Tablets and Starter Kit*)

VENETOCLAX WITH OBINUTUZUMAB FOR PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)

- In combination with obinutuzumab for the treatment of adult patients with previously untreated chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) who are fludarabine ineligible.

Clinical Notes:

- Treatment should be given for a total of 12 months (six 28-day cycles in combination with obinutuzumab, followed by six months of monotherapy), or until disease progression or unacceptable toxicity, whichever occurs first.
- Retreatment with a venetoclax based regimen is funded if relapse is greater than 12 months from completion of venetoclax in combination with obinutuzumab.
- Either ibrutinib or acalabrutinib is funded as a subsequent treatment option, provided all other funding criteria are met.

VENETOCLAX WITH AZACITIDINE FOR NEWLY DIAGNOSED ACUTE MYELOID LEUKEMIA (AML)

- In combination with azacitidine for the treatment of patients with newly diagnosed acute myeloid leukemia (AML) who are 75 years of age or older, or who have comorbidities that preclude the use of intensive induction chemotherapy.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

Clinical Notes:

- Treatment should continue until disease progression or unacceptable toxicity.
- All newly diagnosed AML patients who are ineligible for induction chemotherapy are eligible regardless of cytogenetic risk.,
- On a time-limited need, patients who are currently receiving azacitidine for newly diagnosed AML may have venetoclax added to their treatment provided there is no disease progression and patient otherwise meets criteria.

Claim Notes:

- Patients who have been previously treated with a hypomethylating agent or chemotherapy for the treatment of myelodysplastic syndromes (MDS) are not eligible for treatment with venetoclax in combination with azacitidine.
- Patients with high risk MDS are not eligible for treatment with venetoclax in combination with azacitidine.

VENETOCLAX MONOTHERAPY FOR CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) IN PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

- As a single agent treatment option for patients with chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) who have received at least one prior therapy, and who have failed a B-cell receptor inhibitor (BCRi). Treatment should be continued until disease progression or unacceptable toxicity.

Clinical Note:

- Patients who have an intolerance or a contraindication to a B-cell receptor inhibitor (BCRi) will be eligible for treatment with venetoclax. Intolerance to BCRi would be determined by the clinician.

VENETOCLAX WITH RITUXIMAB FOR CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL) IN PATIENTS WHO HAVE RECEIVED AT LEAST ONE PRIOR THERAPY

- In combination with rituximab for the treatment of adult patients with chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL) who have received at least one prior therapy, irrespective of their 17p deletion status. Treatment should be continued until disease progression or unacceptable toxicity up to a maximum of two years, whichever comes first.

Clinical Notes:

- Patients who were previously treated with an anti-CD20 therapy (rituximab or obinutuzumab) will be eligible if they had a progression-free interval of 6 months or longer. For patients previously treated with venetoclax, the progression-free interval must be 12 months or longer.
- Patients currently receiving and responding to venetoclax monotherapy, and who have not achieved an adequate response are eligible to have rituximab added to venetoclax. Note: Venetoclax therapy is funded to a maximum of two years from the time rituximab is added.
- Patients will be eligible for treatment with either ibrutinib, or idelalisib with rituximab following progression on venetoclax with rituximab if they have not received before and otherwise meet eligibility criteria.

VERTEPORFIN (*Visudyne 15mg/Vial Injection*)

- For the treatment of wet age-related macular degeneration (AMD) as prescribed by an authorized ophthalmologist [Criteria Code 01]

VIGABATRIN (*Sabril 0.5g Sachet and 500mg Tablet*)

- For the treatment of epilepsy in those patients who respond inadequately to alternative treatment combinations, or in whom other drug combinations have not been tolerated, and in whom the potential benefits conferred by its use

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outweigh the risk of ophthalmologic abnormalities.

- For the management of infantile spasms.

***VISMODEGIB** (*Erivedge 150mg Capsule*)

- As a single agent for the treatment of measurable metastatic BCC, OR
- For the treatment of locally advanced BCC (including basal cell nevus syndrome i.e. Gorlin syndrome who are 18 years of age and older) in patients who are inappropriate for surgery and radiotherapy based on a discussion/evaluation with other members of the multi-disciplinary team.
- Patient has ECOG \leq 2

***VITAMIN B₁₂, INJECTION**

- See [Cyanocobalamin, Injection](#)

***VITAMIN B₁₂, ORAL**

- See [Cyanocobalamin, Oral](#)

VORICONAZOLE (*Vfend 50mg, 200mg Tablet and generic brands*)

- For the management of invasive aspergillosis
- For the treatment of culture proven invasive candidiasis with documented resistance to fluconazole

Claim Notes:

- Must be prescribed by a hematologist or specialist in infectious diseases or medical microbiology.
- Initial requests will be approved for a maximum of 3 months.

***WET NEBULIZATION SOLUTIONS** (*Budesonide, Fenoterol, Ipratropium Bromide, Salbutamol*)

- For adult patients with a vital capacity of 900mL or less
- For adult patients with a respiratory rate greater than 25 breaths/minute
- For patients who have demonstrated they cannot follow instructions, cannot hold the spacer device or cannot hold the device long enough to actuate it
- Other requests reviewed on a case by case basis

***ZANAMIVIR** (*Relenza 5mg Powder For Inhalation*)

- For the treatment of long-term care residents with clinically suspected or lab confirmed influenza A or B, when there is documented resistance to oseltamivir or when oseltamivir is contraindicated. A clinically suspected case is one in which the patient meets the criteria of influenza-like illness and there is confirmation of influenza A or B circulating within the facility or surrounding community.
- For the prophylaxis of influenza A or B in long term care residents where the facility has an outbreak, when there is documented resistance to oseltamivir or when oseltamivir is contraindicated.
- A protocol has been developed by Public Health for the treatment of patients in long-term care facilities. The facility must contact the Medical Officer of Health or local Public Health Office, who will notify the Pharmacare office (or dispensing pharmacy after office hours) if coverage is required.

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).

***ZANUBRUTINIB** (*Brukinsa 80mg Capsule*)

PREVIOUSLY UNTREATED CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)

- As monotherapy for the treatment of adult patients with previously untreated chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) for whom a fludarabine-based regimen is considered inappropriate due to a high risk of relapse or refractory disease based on prognostic biomarkers.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be continued until disease progression or unacceptable toxicity.
- High risk for relapse or refractory disease includes 17p deletion, TP53 mutation, 11q deletion and unmutated IGHV.

Claim Notes:

- Patients are not eligible if they have prolymphocytic leukemia or Richter's transformation.
- Requests will not be considered for patients who experience disease progression on a Bruton's tyrosine kinase (BTK) inhibitor or idelalisib.
- Requests will be considered for patients who are not suitable candidates for intravenous therapy.
- Venetoclax with or without rituximab is funded as a subsequent line of therapy in patients who have experienced disease progression during first-line zanubrutinib treatment, provided all other funding eligibility criteria are met.

RELAPSED/REFRACTORY CHRONIC LYMPHOCYTIC LEUKEMIA (CLL) OR SMALL LYMPHOCYTIC LYMPHOMA (SLL)

- As monotherapy for the treatment of adult patients with relapsed or refractory chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) who have received at least one prior systemic therapy.

Clinical Notes:

- Patients should have a good performance status.
- Treatment should be continued until disease progression or unacceptable toxicity.

Claim Notes:

- Patients are not eligible if they have prolymphocytic leukemia or Richter's transformation.
- Requests will not be considered for patients who experience disease progression on a Bruton's tyrosine kinase (BTK) inhibitor or idelalisib.

RELAPSED OR REFRACTORY WALDENSTRÖM MACROGLOBULINEMIA

- For the treatment of adult patients with relapsed or refractory Waldenström macroglobulinemia who have received at least one prior therapy and have not experienced disease progression on a Bruton's tyrosine kinase inhibitor.

Clinical Notes:

- Patients should have a good performance status and no evidence of disease transformation.
- Treatment should be discontinued upon disease progression or unacceptable toxicity.

ZIPRASIDONE (*Zeldox 20mg, 40mg, 60mg, 80mg Capsule and generic brands*)

- For the treatment of schizophrenia and related psychotic disorders (not dementia related) in patients with a history of failure, intolerance, or contraindication to at least one less expensive antipsychotic agent

NOTE: Exception status drugs for Drug Assistance for Cancer Patients are indicated by an asterisk (*).