Exceptional Access Program
Reimbursement Criteria for Frequently Requested Drugs

January 1, 2022
Disclaimer

The information in this document is updated on a regular basis. Although we strive to ensure that all information is accurate at the time of posting, please be aware that some items may be subject to change from time-to-time.

The following list of drugs and indications that will be considered for funding under the Exceptional Access Program is not exhaustive. Physicians may wish to contact the EAP directly by phone at 416-327-8109 or 1-866-811-9893 or by email at EAPFeedback.MOH@ontario.ca to see if a specific drug product and or indication not listed below may be considered for EAP funding.

The information provided in this document and website is intended for information purposes only and does not provide any medical diagnosis, symptom assessment, health counselling or medical opinion for individual users. This information also does not constitute medical advice for physicians or patients. For more detailed information on prescription drugs, please consult a qualified healthcare professional.
Reimbursement Criteria for Frequently Requested Drugs and Indications

For a drug to be considered for funding, the EAP reimbursement criteria must always be met and the request approved prior to the initiation of treatment with the drug being requested, unless otherwise specified within the criteria. This includes:

- funding for continued treatment that was previously supplied through a clinical trial, or paid for by other means (such as a third party payer)
  
  *Note:* First time applications for the funding of ongoing treatments must meet both initial and renewal criteria for the drug being requested (unless otherwise specified)

- funding for a renewal beyond the previously approved initial period, unless otherwise specified.

Note that the terms “fund”, “funded”, or “funding” within this document are interpreted and applied by the Ministry in accordance with the clinical evidence used to establish the reimbursement criteria. The Ministry does not distinguish between the source of the drug funding (e.g. public or private payer[s]) in administering the EAP reimbursement criteria.

Consider the following:

*The “non-funding” of specific combination treatments as identified by the criteria will not be reimbursed regardless of the funding source(s) of either therapy. The funding criteria are established based on the reviewed clinical evidence through the submission process and are not to be misconstrued with the source(s) of funding.*

Example 1: If drug A has approved reimbursement criteria and drug B has approved reimbursement criteria but the combination therapy of drug A plus drug B has not been reviewed through the established process and/or has no reimbursement criteria, the EAP will not fund either drug individually or as combination therapy if the intended use is for combination therapy, regardless of the actual source of funding of either drug A or drug B.

Example 2: If drug A has approved reimbursement criteria and drug B does not have approved reimbursement criteria, and drug A used in combination with drug B does not have approved reimbursement criteria, funding for drug A will not be provided if the intent is to use drug A as combination with drug B, regardless of the source of funding for drug B.

*The duration of funding of a regimen identified in the criteria is in accordance with the duration of therapy supported by the clinical evidence and is not related to or dependent on source(s) of funding.*

Example 1: If the approved reimbursement criteria states that “The Ministry will fund drug C for a period of 3 years” and drug C was already used by the patient for 2 years funded by another payer (e.g. private payer, manufacturer, out-of-pocket), the Ministry will only be obliged to fund drug C for the remaining one year if the request meets the approved EAP reimbursement criteria. Such a limitation in the duration of funding is aligned with the clinical evidence provided to the Ministry at the time of the review.
For a limited number of requests where expert opinion is required, the requests are reviewed by an external reviewer who is a medical expert in the field.

Where available, a link has been provided to the information page containing details of the Committee to Evaluate Drugs (CED) review and subsequent the Executive Officer’s funding decision for the particular drug and indication. Information on whether the drug and indication can be considered through the Telephone Request Service (TRS) is also included.

EAP requests may be submitted for numerous other drugs not listed below, or for drugs listed below but for different indications. However, EAP funding will only be considered for drugs and indications that have been reviewed by the CED and approved for funding by the Executive Officer. For more information, please refer to the main EAP webpage.

Some of the drugs considered through EAP are also listed on the ODB Formulary for a different indication as Limited Use (LU) benefit. You can check whether the drug is listed by searching the e-Formulary.

For details on how the EAP reimbursement criteria are developed, please refer to the main EAP webpage.

To assist physicians applying for exceptional access, the ministry has developed a standard form.

Use of form is not mandatory but does facilitate provision of all relevant information. Where applicable, please ensure that all relevant clinical information is provided demonstrating that the patient meets the reimbursement criteria.

*Note: The dosage form and strength of the product that has been approved for reimbursement consideration are those that have been approved by the Committee to Evaluate Drugs (CED). In most cases, these are the dosage forms and strengths submitted to the CED by the manufacturer for consideration, however, it may not be inclusive of all dosage forms and strengths available through the manufacturer.*
**Biosimilar Policy and the Reimbursement of Biologic Originators through The Exceptional Access Program**

Reimbursement for Remicade®, Enbrel®, Humira®, Rituxan®, and Copaxone® (not a biologic but a complex molecule included under this policy) through the Ontario Drug Benefit (ODB) program is provided only for eligible ODB recipients who are treatment experienced on a case-by-case basis. All patients must meet specified initiation criteria and renewal criteria associated with the treatment depending on the duration of use of the biologic by the patient. This requirement must be met irrespective of whether the drug was originally funded through a private health insurance plan, a clinical trial, compassionately by another payer, accessed in another country or jurisdiction, or paid for out-of-pocket and it is intended to optimize consistency and equity of funding for Ontarians in accordance with the evidence-review processes.

Patients who are treatment naïve to the biologic are funded for the biosimilar version for its Health Canada approved indications. The biosimilar version of these drugs can be accessed as a limited use benefit upon meeting specified criteria for their Health Canada approved indications. If a request does not meet the LU criteria on the ODB formulary, case-by-case consideration may be provided through the EAP.

Patients must be provided with a new prescription and meet the Limited Use Criteria or Exceptional Access Program criteria to be publicly reimbursed for either the biologic originator or its biosimilar versions. The biosimilar version is generally available for all its Health Canada approved indications on the ODB formulary as a limited use benefit. Additionally, requests for a biologic for indications where Health Canada has not granted notice of compliance for the biologic originator or the biosimilar versions, if approved for funding through a case-by-case process, will be granted access to the biologic originator and the biosimilar versions.

Please visit the Ministry website to view the more detailed information pertaining to each biologic originator/biosimilar announcement in the Executive Officer Communication Notices. Frequently Asked Questions pertaining to each biologic/drug at the time of implementation are also found on the Ministry website.


You can find the limited use criteria associated with funded biosimilars on the e-formulary or the latest version of the Ontario Drug Benefit Formulary. The searchable electronic formulary is available at the following URL on the Drugs and Devices Division’s website:

https://www.formulary.health.gov.on.ca/formulary/
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ANEMIA

Darbepoetin
Brand(s): Aranesp
DOSAGE FORM/ STRENGTH: Prefilled syringes: 150 mcg, 200 mcg, 300 mcg, 500 mcg

(Requests for the treatment of chemotherapy-induced anemia in patients with malignant cancer DO NOT require an EAP submission. Please refer to the ODB e-formulary to determine if the patient satisfies the criteria for use.)

For the treatment of anemia secondary to chronic renal disease in those who are not eligible under the Special Drugs Program, approval can be given if the patient meets the following criteria:

Estimated glomerular filtration rate (GFR) less than 30 mL/min AND
Baseline hemoglobin level less than 100 g/L AND
Mean corpuscular volume (MCV) level between 75 fL and 120 fL

All requests MUST indicate the reason why the patient is ineligible for the Special Drugs Program.

Duration of Approval: 6 months

Renewals will be provided to patients where the hemoglobin levels have improved by 15 g/L after 3 months of therapy.

Renewals must specify the name of the drug and dose requested and MUST be accompanied by bloodwork that includes a recent hemoglobin level. Also, please identify if the patient has received transfusions after the first 2 weeks of therapy with darbepoetin and the date(s) that the transfusion(s) occurred.

Duration of Approval: 12 months

For the treatment of anemia secondary to myelodysplastic syndrome (MDS) in patients who meet the following criteria:

- MDS confirmed by the bone marrow report AND
- With a hemoglobin count less than 100 g/L AND
- Endogenous erythropoietin level of less than 500 U/L AND
- Mean corpuscular volume (MCV) level between 75 fL and 120 fL.

Submissions must include the date(s) for the above blood work.
Darbepoetin
Brand(s): Aranesp
DOSAGE FORM/ STRENGTH: Prefilled syringes: 150 mcg, 200 mcg, 300 mcg, 500 mcg

For patients with an MCV level below 75 fL or above 120 fL, the physician must provide a discussion of how reversible causes of anemia were ruled out to enable further consideration of the submission.

Duration of Approval: 6 months

Renewals will be provided to patients where the hemoglobin levels have improved by 15 g/L after 3 months of therapy.

Renewals must specify the name of the drug and dose requested and MUST be accompanied by bloodwork that includes a recent hemoglobin level. Also, please identify if the patient has received transfusions after the first 2 weeks of therapy with darbepoetin and the date(s) that the transfusion(s) occurred.

Duration of Approval: 12 months

For the treatment of anemia secondary to myelodysplastic syndrome (MDS) in patients who meet the following criteria:

- MDS confirmed by the bone marrow report AND
- With a hemoglobin count less than 100 g/L AND
- Endogenous erythropoietin level of less than 500 U/L AND
- Mean corpuscular volume (MCV) level between 75 fL and 120 fL.

Submissions must include the date(s) for the above blood work.

For patients with an MCV level below 75 fL or above 120 fL, the physician must provide a discussion of how reversible causes of anemia were ruled out to enable further consideration of the submission.

Duration of Approval: 6 months

Renewals will be provided to patients where the hemoglobin levels have improved by 15 g/L after 3 months of therapy.

Renewals must specify the name of the drug and dose requested and MUST be accompanied by bloodwork that includes a recent hemoglobin level. Also, please identify if the patient has received transfusions after the first 2 weeks of therapy with darbepoetin and the date(s) that the transfusion(s) occurred.

Duration of Approval: 12 months
**Epoetin Alpha**

**Brand(s): Eprex**

**DOSAGE FORM/ STRENGTH:** Prefilled syringes: 1,000 IU, 2,000 IU, 3,000 IU, 4,000 IU, 5,000 IU per 0.5 mL, 6,000 IU/0.6 mL, 8,000 IU/0.8 mL, 10,000 IU/mL, 20,000 IU/0.5 mL, 40,000 IU/mL; Check the formulary and/or e-formulary for funded products

For the treatment of anemia secondary to chronic renal disease in those who are not eligible under the Special Drugs Program, approval can be given if the patient meets the following criteria:

- Estimated glomerular filtration rate (GFR) less than 30 mL/min **AND**
- Baseline hemoglobin level less than 100 g/L **AND**
- Mean corpuscular volume (MCV) level between 75 fL and 120 fL

All requests MUST indicate the reason why the patient is ineligible for the Special Drugs Program.

**Renewals** will be provided to patients where the hemoglobin levels have improved by 15 g/L after 3 months of therapy.

Renewals must specify the name of the drug and dose requested and MUST be accompanied by bloodwork that includes a recent hemoglobin level. Also, please identify if the patient has received transfusions after the first 2 weeks of therapy with epoetin alpha and the date(s) that the transfusion(s) occurred.

For the treatment of anemia secondary to myelodysplastic syndrome (MDS) in patients who meet the following criteria:

- MDS confirmed by the bone marrow report **AND**
- With a hemoglobin count less than 100 g/L **AND**
- Endogenous erythropoietin level of less than 500 U/L **AND**

Mean corpuscular volume (MCV) level between 75 fL and 120 fL.

Submissions must include the date(s) for the above blood work.

For patients with an MCV level below 75 fL or above 120 fL, the physician must provide a discussion of how reversible causes of anemia were ruled out to enable further consideration of the submission.

**Duration of Approval:** 6 months

**Renewals** will be provided to patients where the hemoglobin levels have improved by 15 g/L after 3 months of therapy.
**Epoetin Alpha**
Brand(s): Eprex
**DOSAGE FORM/ STRENGTH:** Prefilled syringes: 1,000 IU, 2,000 IU, 3,000 IU, 4,000 IU, 5,000 IU per 0.5mL; 6,000 IU/0.6 mL; 8,000 IU/0.8 mL; 10,000 IU/mL; 20,000 IU/0.5 mL, 40,000 IU/mL

Renewals must specify the name of the drug and dose requested and MUST be accompanied by bloodwork that includes a recent hemoglobin level. Also, please identify if the patient has received transfusions after the first 2 weeks of therapy with epoetin alfa and the date(s) that the transfusion(s) occurred.

**Duration of Approval:** 6 months

**For treatment of anemia secondary to hepatitis C therapy** in patients who meet the following criteria:

- Patient is diagnosed with Hepatitis C and is undergoing treatment with pegylated interferon and ribavirin **AND**
- Current hemoglobin is >100 g/L but patient has experienced a drop in hemoglobin of at least 40 g/L since treatment **OR**
- Current hemoglobin count is < 100 g/L

Submissions must include the following:

1) Details of therapy with pegylated interferon and ribavirin (ie: Start date, duration of treatment, patient response etc)

2) Bloodwork (full CBC preferred) that includes the hemoglobin and mean corpuscular volume (MCV) as well as the date(s) for the above blood work.

For patients with an MCV level below 75 fL or above 120 fL, the physician must provide a discussion of how reversible causes of anemia were ruled out to enable further consideration of the submission.

Submissions not meeting the above criteria will be considered on a case-by-case basis. All submissions should be accompanied by

- Baseline and current bloodwork (full CBC with MCV)
- Baseline clinical status and current symptoms from anemia that were not present at baseline
- Details of any complications from anemia

**Duration of Approval:** Full duration of treatment with pegylated interferon/ribavirin treatment
Epoetin Alpha
Brand(s): Eprex
DOSAGE FORM/ STRENGTH: Prefilled syringes: 1,000 IU, 2,000 IU, 3,000 IU, 4,000 IU, 5,000 IU per 0.5mL; 6,000 IU/0.6 mL; 8,000 IU/0.8 mL; 10,000 IU/mL; 20,000 IU/0.5 mL, 40,000 IU/mL

Renewals will be granted for the full period of pegylated interferon and ribavirin treatment in those who show significant response to therapy. Renewals should be accompanied by bloodwork that includes a recent hemoglobin and must identify if the patient has required transfusions after the first 2 weeks of therapy.

Duration of Approval: Full duration of treatment with pegylated interferon/ribavirin treatment

Pre-operative use at a dose up to 40,000 IU weekly prior to single hip, double knee, or single (“redo”) knee surgery in patients who meet the following criteria;

- Hemoglobin between 100 – 130 g/L inclusive AND
- Mean corpuscular volume (MCV) level between 75 fL and 120 fL inclusive

Request not meeting these criteria will be assessed on a case-by-case basis.

Duration of Approval: Up to 4 doses preoperatively

For the treatment of anemia in palliative cancer patients. Individuals will be assessed on a case-by-case basis. Submissions must include the rationale for using epoetin alpha over transfusion.

Requests for the treatment of chemotherapy-induced anemia in patients with malignant cancer DO NOT require an EAP submission. Please refer to the e-formulary to determine if the patient satisfies the criteria for use.
Iron dextran complex
Brand(s): Dexiron
DOSAGE FORM/ STRENGTH: 50 mg/mL Injectable
Product is dormant but patients with an existing EAP approval may use remaining supplies as appropriate

For the treatment of iron-deficiency anemia confirmed by bloodwork where the patient has a demonstrated intolerance\(^1\) to oral iron therapy\(^2\) OR has not responded to adequate therapy with oral iron\(^2\).

\(^1\)Intolerance must be described.

\(^2\)Provide name of iron salt, dose, duration of therapy, response etc.

Duration of Approval: 1 year
Renewals will be considered on a case-by-case basis.

Duration of Approval: 2 years

Iron sucrose
Brand(s): Venofer, PMS Iron Sucrose
DOSAGE FORM/ STRENGTH: 20 mg/mL Injectable

For the treatment of iron-deficiency anemia confirmed by bloodwork where the patient has a demonstrated intolerance\(^1\) to oral iron therapy\(^2\) OR has not responded to adequate therapy with oral iron\(^2\).

\(^1\)Intolerance must be described.

\(^2\)Provide name of iron salt, dose, duration of therapy, response etc.

Duration of Approval: 1 year
Renewals will be considered on a case-by-case basis.

Duration of Approval: 2 years

Iron isomaltoside 1000
Brand(s): Monoferric
DOSAGE FORM/ STRENGTH: 100 mg/mL Injectable

Requests for iron isomaltoside 1000 (Monoferric) 100 mg/mL that do not meet the Limited Use Criteria on the Ontario Drug Benefit formulary may be considered for funding on a case-by-case basis by the EAP.
Lenalidomide
Brand(s): Revlimid
DOSAGE FORM/STRENGTH: 2.5 mg, 5 mg, 10 mg, 15 mg, 20 mg, 25 mg capsule

For the treatment of anemia due to myelodysplastic syndrome (MDS) for patients who meet all the following clinical criteria;

- Demonstrated diagnosis of MDS on bone marrow aspiration; AND
- Presence of del[5q] documented by standard cytogenetic or fluorescence in situ hybridization; AND
- International Prognostic Scoring System (IPSS) risk category low or intermediate-1; AND
- Transfusion-dependent symptomatic anemia.

Duration of Approval: 6 Months

Renewal will be considered for patients who are transfusion-dependent and who have demonstrated at least a fifty percent (50%) reduction in transfusion requirements.

Duration of Approval: Up to 1 Year

Note: Patients with anemia due to MDS who are not transfusion-dependent will be assessed on a case-by-case basis.

- Physicians submitting initial requests for non-transfusion-dependent patients must provide clinical evidence of symptomatic anemia affecting the patient’s quality of life and the rationale for why transfusions are not being used.
- Renewal requests for non-transfusion-dependent patients will be considered on a case-by-case basis. In such cases, the requesting physician will be required to provide the patient’s serial clinical blood culture (CBC) pre- and post-lenalidomide therapy in addition to any other objective evidence of the patient’s response to lenalidomide therapy.
**ANTICOAGULANTS**

**Dalteparin**
Brand(s): Fragmin

**DOSAGE FORM/ STRENGTH:** Check formulary or e-formulary for funded products

For **peri-operative bridging** for patients who require long-term warfarin therapy and must temporarily discontinue it before and after surgery, and who are at moderate- to high-risk for an embolic event while off warfarin.

**Standard Approval Duration:** As requested up to a maximum of **10 days** before the date of surgery **plus up to 7 days** after surgery.

For **post-operative prophylaxis of DVT** for patients who had hip or knee surgery, and cannot use warfarin.

**Standard Approval Duration:** As requested up to a maximum of **30 days** starting on the day of surgery.

For the **post-operative prophylaxis of venous thromboembolism following abdominal or pelvic surgery for cancer in patients** who do not have a history of or risk factors for heparin-induced thrombocytopenia.

**Standard Approval Duration:** Maximum of 30 days

For **extended treatment of symptomatic acute venous thromboembolism (VTE)** in patients with cancer for whom warfarin treatment is not appropriate on a case-by-case basis.

**Standard Approval Duration:** 6 months

**Enoxaparin – See formulary for funded biosimilars**
Brand(s): Lovenox and biosimilars

**DOSAGE FORM/ STRENGTH:** Check formulary or e-formulary for funded products

**Tinzaparin**
Brand(s): Innohep

**DOSAGE FORM/ STRENGTH:** Check formulary or e-formulary for funded products

For **peri-operative bridging** for patients who require long-term warfarin therapy and must temporarily discontinue it before and after surgery, and who are at moderate- to high-risk for an embolic event while off warfarin.

**Standard Approval Duration:** As requested up to a maximum of **10 days** before the date of surgery **plus up to 7 days** after surgery.
**Enoxaparin**
Brand(s): Lovenox
DOSAGE FORM/ STRENGTH: Check formulary or e-formulary for funded products

**Tinzaparin**
Brand(s): Innohep
DOSAGE FORM/ STRENGTH: Check formulary or e-formulary for funded products

For **post-operative prophylaxis of DVT** for patients who had hip or knee surgery, and cannot use warfarin.

**Standard Approval Duration:** As requested up to a maximum of **30 days** starting on the day of surgery.

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For the **post-operative prophylaxis of venous thromboembolism following abdominal or pelvic surgery for cancer in patients** who do not have a history of or risk factors for heparin-induced thrombocytopenia.

Standard Approval Duration: Maximum of 30 days

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**Fondaparinux**
Brand(s): Arixtra and generic
DOSAGE FORM/ STRENGTH: 2.5 mg

For the **post-operative prophylaxis of venous thromboembolism following abdominal or pelvic surgery for cancer**.

**Standard Approval Duration:** Maximum of 30 days

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**Fondaparinux**
Brand(s): Arixtra (2.5mg) and Generic (2.5mg and 7.5mg) Check formulary or e-formulary for funded products
DOSAGE FORM/ STRENGTH: 2.5 mg and 7.5 mg

For the treatment of **venous thromboembolism (VTE)** [deep vein thrombosis (DVT), pulmonary embolism (PE)] in the setting of acute heparin induced thrombocytopenia (HIT)

Duration: 1 month if patient is able to transition to warfarin OR up to 1 year for patients who cannot be managed on warfarin*
**Fondaparinux**

Brand(s): Arixtra (2.5mg) and Generic (2.5mg and 7.5mg) Check formulary or e-formulary for funded products

**DOSAGE FORM/ STRENGTH:** 2.5mg and 7mg Injection

For the treatment of venous thromboembolism (VTE) [deep vein thrombosis (DVT), pulmonary embolism (PE)] in patients with previous HIT who cannot use rivaroxaban

Duration: 1 month if patient is able to transition to warfarin OR up to 1 year for patients who cannot be managed on warfarin*

* Patients may not be able to use warfarin as extended treatment or prophylaxis of VTE due to contraindication, previous clinical failure, inability to swallow/ take oral medications, immobility, inability to monitor INR, etc. The reasons why warfarin is not appropriate must be clearly described.

Renewal of funding for the above indications may be considered on a case-by-case basis with documented clinical rationale that includes details pertaining to patient’s risks for recurrent VTE, risks of bleeding, update on whether warfarin therapy may now be considered, and response to therapy.

For peri-operative bridging for patients with history of HIT who require long-term warfarin therapy and must temporarily discontinue it before and after surgery, and who are at moderate- to high-risk for an embolic event while off warfarin.

Duration: As requested up to a maximum of 10 days before the date of surgery plus up to 7 days after surgery

Renewal of funding will NOT be considered for this indication.

**Nadroparin**

Brand(s): Fraxiparine

**DOSAGE FORM/ STRENGTH:** Check formulary or e-formulary for funded products

For the post-operative prophylaxis of venous thromboembolism following abdominal or pelvic surgery for cancer in patients who do not have a history of or risk factors for heparin-induced thrombocytopenia.

Standard Approval Duration: Maximum of 30 days
# ANTICONVULSANTS

## Lamotrigine (chewable)
**Brand(s):** Lamictal  
**DOSAGE FORM/ STRENGTH:** 5 mg chewable tablet

For the adjunctive therapy for children over 2 years of age who are suffering from refractory seizures associated with Lennox-Gastaut syndrome, and who have previously tried other antiepileptic drugs.

**Duration of Approval:** 1 year

## Oxcarbazepine
**Brand(s):** Trileptal  
**DOSAGE FORM/ STRENGTH:** 150 mg, 300 mg, and 600 mg tablet 60 mg/mL

For the treatment of partial seizures in adults and in children aged 6 years and older who have had an inadequate response or intolerance* to at least 3 other formulary agents (prior or current use) including carbamazepine.

* Intolerance must be described in detail.

**Warning:** Life-threatening dermatological reactions, including Stevens Johnson Syndrome and toxic epidermal necrolysis, and multi-organ hypersensitivity reactions have been associated with the use of oxcarbazepine. More information may be found on the Health Canada webpage:  

**Duration of Approval:** Lifetime

## Rufinamide
**Brand(s):** Banzel  
**DOSAGE FORM/ STRENGTH:** 100 mg, 200 mg, 400 mg

For the treatment of seizures associated with Lennox-Gastaut Syndrome (LGS) in patients who meet the following criteria:

- Patient is 4 years of age or older; AND
- the Patient is currently on two or more anti-epileptic drugs (AEDs) without optimal seizure control; AND
- the Patient has failed an adequate trial\(^1\) of lamotrigine AND topiramate; AND
- the Patient is in the care of a physician experienced in managing seizures.
If an adequate trial of lamotrigine and/or topiramate is not possible due to intolerance or contraindication, a less costly AED that is listed as a benefit on the Ontario drug benefit formulary must be tried in its place.

**Dose:** Maximum daily dose is 1,300 mg per day for patients less than 30 kg; and 3,200 mg per day for patients 30 kg or greater.

**Exclusion Criteria.**

Funding will not be approved for the following circumstances:

- Banzel used first line for LGS; OR

Treatment of partial seizures

**Duration of Approval:** Lifetime

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**Stiripentol**

*Brand(s): Diacomit*

*DOSAGE FORM/ STRENGTH:* 250 mg capsule, 250 mg/pack powder for suspension, 500 mg capsule, 500 mg/pack powder for suspension

For the treatment of patients with severe myoclonic epilepsy in infancy (Dravet syndrome) who meet the following criteria;

1. the patient has refractory generalized tonic-clonic seizures; AND
2. the patient requires Diacomit (Stiripentol) for use in combination with clobazam and valproate as adjunctive therapy for the seizures; AND
3. the patient’s seizures are not adequately controlled with clobazam and valproate alone; AND
4. the request is submitted by a neurologist or pediatrician.

Case-by-case consideration through external review will be permitted for circumstances not meeting the above criteria.

**Duration of Approval:** Lifetime
**ANTIDIABETIC AGENTS**

**Pioglitazone**  
*Brand(s):* Actos, Generics  
**DOSAGE FORM/ STRENGTH:** 15 mg, 30 mg, 45 mg tablet

For the treatment of type 2 diabetes in patients who require;

- Dual combination therapy of diabetes AND demonstrate inadequate glycemic control (HbA1c of >7%) on maximal doses of metformin (2000 mg/day) OR
- Dual combination therapy of diabetes AND demonstrate inadequate glycemic control (HbA1c of >7%) on maximal* doses of sulfonylurea and demonstrated intolerance / contraindication to metformin OR
- Triple combination therapy of diabetes and who demonstrate inadequate glycemic control on maximal** doses of metformin and a sulfonylurea AND only if the physician has offered insulin as an alternative option first, and the patient has refused or is not able to take insulin. Note: Both the physician and patient must be aware that thiazolidinediones (TZDs), are not indicated for use in triple therapy.

***Those with one or more of the following contraindications/ precautions to therapy with pioglitazone/rosiglitazone will not be considered

- Patients with type 1 diabetes
- Patients who will be using this as monotherapy
- Combination use with a nitrates
- Combination use with insulin
- Patients with any stage of heart failure (i.e. NYHA Class I, II, III, IV)
- Patients at high risk for bone fracture (i.e. post-menopausal women with previously confirmed osteoporosis or osteopenia)
- Patients with recent history (in the past 3 months) of an ischemic cardiovascular event (myocardial infarction, unstable angina)

* Note: For the purpose of the EAP submission, maximal dose of sulfonylurea is considered to be glyburide 10 mg/day, gliclazide 160mg/day OR Diamicron MR 60mg/day, OR glimepiride (Amaryl ) 4 mg/day and aximal dose of metformin is considered to be 2000 mg/day.

**Duration of Approval:** 5 years

**Renewals** as well as requests for ongoing treatment in patients previously provided these drugs by other means will be considered for those patients who have NOT developed a contraindication/precautionary use*** in the intervening period AND have demonstrated a recent HbA1c level ≤7% while on treatment

**Duration of Approval:** 5 years
Repaglinide  
Brand(s): GlucoNorm  
DOSAGE FORM/ STRENGTH: 0.5 mg, 1 mg, 2 mg tablet

For the treatment of type 2 diabetes in patients with:

- Inadequate glycemic control (HbA1c >7%) using maximal* doses of a sulfonylurea AND metformin (2000mg/day) OR

- Inadequate glycemic control and demonstrated intolerance or contraindication to metformin and who are on maximal* doses of a sulfonylurea OR

- Inadequate glycemic control and demonstrated intolerance or contraindication to a sulfonylurea (glyburide, gliclazide or glimepiride) and are on maximal** doses of metformin OR

- Demonstrated intolerance or contraindication to both a sulfonylurea (glyburide, gliclazide or glimepiride) AND metformin OR

- Adequate glycemic control (HbA1c ≤ 7%) who develops intolerance or contraindication to sulfonylurea (glyburide, gliclazide or glimepiride) or metformin OR

- HbA1c ≤ 7% but with greater than 50% of fasting blood glucose (FBG >7mmol/L) or post-prandial plasma glucose (PPG >10mmol/L) levels not within target range and using maximally tolerated doses of a sulfonylurea and metformin.

* Note: For the purpose of the EAP submission, maximal dose of sulfonylurea is considered to be glyburide 10mg/day, gliclazide 160 mg/day or Diamicron MR 60 mg/day, OR glimepiride (Amaryl) 4 mg/day.

**Note: For the purpose of the EAP submission, maximal dose of metformin is considered to be 2000 mg/day

Duration of Approval: 5 years
Rosiglitazone
Brand(s): Avandia
DOSAGE FORM/ STRENGTH: 2 mg, 4 mg, 8 mg tablet

For the treatment of type 2 diabetes mellitus in patients with:

- Inadequate glycemic control (HbA1c >7%) from ALL other oral antidiabetic agents* funded through one of the Ontario Drug Benefit Programs, in monotherapy or in combination OR

- Where ALL other oral antidiabetic agents are inappropriate due to contraindications or intolerance AND

- The patient has refused or is not able to take insulin AND

- There is no known contraindication to rosiglitazone

* Oral antidiabetics include the following agents;
  - glyburide
  - metformin
  - gliclazide (Diamicron, Diamicron MR)
  - sitagliptin (Januvia)
  - saxagliptin (Onglyza)
  - repaglinide (GlucoNorm)
  - pioglitazone (Actos)

Note: A trial with acarbose is not a mandatory requirement.

Note: It is not necessary for patients to have tried the following oral antidiabetic agents that are currently not funded by the OPDP for the purposes of obtaining rosiglitazone:
  - glimepiride (Amaryl)
  - nateglinide (Starlix)

Duration of Approval: 5 years

Renewals will be considered where patients have benefited and continue to benefit from rosiglitazone treatment as demonstrated by recent HbA1c levels ≤7% while on treatment with rosiglitazone AND in those who continue to have no known contraindication(s) to rosiglitazone.

Duration of Approval: 5 years
Orlistat
Brand(s): Xenical
DOSAGE FORM/ STRENGTH: 120 mg capsule

For the treatment of type 2 diabetes in a patient with:

- Inadequate glycemic control (i.e., HbA1c > 7.0%) on maximal oral antidiabetic medications* AND
- Body Mass Index ≥ 27 AND
- Demonstrated failure to a trial of nutritional/dietary counselling and exercise programs

* Note: Maximal dose of sulfonylurea is considered to be glyburide 10mg/day, gliclazide (160mg/day or Diamicron MR 60 mg/day) OR glimepiride (Amaryl ) 4mg/day.

Note: Maximal dose of metformin is considered to be 2000 mg/day

Duration of Approval: 1 year

Renewals will be considered for those with demonstrated response to treatment reported as at least 5% weight loss and improvement in glycemic control (i.e., HbA1c <7.0% or HbA1c reduction of more than 0.5%)

Duration of Approval: 12 months (First Renewal)
ANTI-INFECTIVES

Aztreonam
Brand(s): Cayston
DOSAGE FORM/ STRENGTH: 75 mg/vial powder for solution

For the treatment of chronic infection with *Pseudomonas aeruginosa* (PsA) infection in patients with a diagnosis of Cystic Fibrosis who meet all the following criteria:

(a) Patient has a documented diagnosis of cystic fibrosis;
(b) Patient has a chronic infection with *Pseudomonas aeruginosa* (PsA) that has been confirmed by 2 (two) positive sputum cultures taken at least 1 month apart that are both positive for PsA;
(c) the Patient’s clinical condition is deteriorating despite treatment with inhaled tobramycin;
(d) the Patient has moderate to severe impairment of lung function defined by baseline FEV1 < 75% of predicted; and
(e) the Patient is ≥ 6 years old.

**Exclusion Criteria:** Aztreonam (Cayston) will not be funded in the following circumstances.
- Aztreonam will not be funded in combination with tobramycin inhalation
- Aztreonam will not be funded for bronchiectasis indications outside of proven cystic fibrosis;
- Aztreonam will not be funded outside of the cystic fibrosis population
- Aztreonam will not be funded for patients with mild cystic fibrosis;
- Aztreonam will not be funded for the purpose of convenience

**Approved Dosage.** The approved dosage for Aztreonam (Cayston) under the EAP is as follows:

Inhale 75 mg three times daily used in a repeated 28 day cycle that involves administration of aztreonam for 4 weeks of treatment followed by 4 weeks off aztreonam therapy.

**Duration of Approval:** 1 year
### Aztreonam
**Brand(s):** Cayston  
**DOSAGE FORM/ STRENGTH:** 75 mg/vial powder for solution

**Renewals** will be considered in patients who demonstrate ongoing response to therapy.

**Duration of Approval:** 1 year

**Approved Dosage.** The approved dosage for Aztreonam (Cayston) under the EAP is as follows:

Inhale 75 mg three times daily used in a repeated 28 day cycle that involves administration of aztreonam for 4 weeks of treatment followed by 4 weeks off aztreonam therapy.

**Renewals** will be considered in patients who demonstrate ongoing response to therapy.

**Duration of Approval:** 1 year

### Dapsone
**Brand(s):** Dapsone  
**DOSAGE FORM/ STRENGTH:** 100 mg tablet

Note that Effective on July 31, 2018, Dapsone has been moved to the Ontario Drug Benefit (ODB) Formulary as a general benefit.

### Daptomycin
**Brand(s):** Cubicin RF and generic daptomycin (Refer to the formulary for funded list)  
**DOSAGE FORM/ STRENGTH:** 500 mg/10 mL powder for injection

Note that initial requests for Daptomycin may be accessed through the EAP’s Telephone Request Service.

**For the treatment of patients experiencing the following types of infections due to methicillin-resistant Staphylococcus aureus (MRSA) bacteria:**

i) Staphylococcus aureus bloodstream (SAB) infection including right-sided Staphylococcus aureus infective endocarditis (SARIE); AND/OR

ii) Osteomyelitis; AND/OR

iii) Device-related osteoarticular or prosthetic joint infections; AND/OR

iv) Diabetic foot infections.

Additionally, the patient must have failed to adequately respond to, be intolerant\(^1\) to, or have a contraindication to vancomycin.
Daptomycin
Brand(s): Cubicin RF, generic daptomycin products (refer to the formulary list)
DOSAGE FORM/ STRENGTH: 500 mg/10 mL powder for injection

1Intolerance due to Red Man Syndrome. If the physician asserts that the patient is intolerant to vancomycin due to red man’s syndrome, additional clinical details of the patient’s intolerance, including rate of infusion and the use of antihistamines and other histamine blockers prior to therapy with vancomycin.

Duration of Approval: Up to 8 weeks

Renewals will be considered on a case-by-case basis. (Physicians must submit adequate clinical information to justify the need for ongoing therapy with daptomycin.)

Duration of Approval: Case-by-case

Exclusion Criteria:

- Daptomycin is not funded for patients with MRSA-related pneumonia;
- Daptomycin is not funded for patients with skin/skin structure infections other than diabetic foot infections caused by MRSA.

For the treatment of invasive infections1 caused by vancomycin-resistant enterococcus (VRE) in patients who meet the following criteria:

1. VRE infection is confirmed by blood or tissue culture and sensitivity report
2. Patient is unable to use linezolid as a result of at least ONE of the following reasons:
   i) has developed resistance to linezolid as confirmed by the microbiology sensitivity report
   ii) has experienced intolerance2 to linezolid (for example: severe gastric symptoms, myelosuppression, peripheral neuropathy requiring medical intervention, lactic acidosis)
   iii) has a contraindication2 to linezolid

OR

Prescribed by an infectious disease expert for a patient who is able to use linezolid but where the bacteriostatic effect of linezolid may not be deemed to be clinically optimal due to other patient factors or comorbidities (e.g., immunocompromised, neutropenic).

1Not approved for colonization (e.g., nares, skin, stool)
Intolerances and contraindications to linezolid must be fully described in the EAP application.

**Daptomycin**
**Brand(s):** Cubicin RF, generic daptomycin (refer to the formulary for funded list)
**DOSAGE FORM/ STRENGTH:** 500 mg/10 mL powder for injection

**Recommended dose:** 8 to 12 mg/kg daily for VRE with adjustments based on renal function

**Duration of approval:**

*Note that the below are examples of durations for reimbursement of some common VRE infections. This is not an all-inclusive list.*

- Urinary tract infections: up to 10 days
- Bacteremia: up to 14 days
- Endocarditis: up to 6 weeks
- Osteomyelitis: up to 8 weeks

Requests for longer durations of funding will be considered case-by-case through external review and must be accompanied by a recent microbiology sensitivity report to confirm sensitivity to daptomycin.
Fidaxomicin (May be accessed through the telephone request service)
Brand(s): Dificid
DOSAGE FORM/ STRENGTH: 200 mg tablet

For the treatment of Clostridium difficile infection (CDI) in patients who meet the EAP criteria for vancomycin use, but 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 failure* 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 intolerance** to oral vancomycin treatment that resulted in the discontinuation of vancomycin therapy.

*Treatment failure is defined as 7 days of vancomycin therapy without acceptable clinical improvement.

**Details of severe adverse reaction or intolerance must be provided and should be clinically related to oral administration of vancomycin.

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: Fecal biotherapy (“stool transplantation”), if available, should be encouraged for this patient population.

Approved dose and duration: 200 mg twice a day for 10 days
Isavuconazole  
Brand(s): Cresemba  
DOSAGE FORM/ STRENGTH: 100 mg capsule, 200 mg Injection Solution

For the treatment of invasive aspergillosis in patients meeting the following criteria:

- 18 years of age and older; AND
- Patient has failed, experienced intolerance to, or has contraindications to voriconazole; AND
- Isavuconazole is prescribed by or in consultation with an infectious disease specialist. (Include consult note with the request)

Exclusion criteria:

Patients with familial short QT syndrome.

Approval Duration: 3 months

Renewals will be considered on a case-by-case basis.

For the treatment of invasive mucormycosis in patients meeting the following criteria:

- 18 years of age and older; AND
- Isavuconazole is prescribed by or in consultation with an infectious disease specialist. (Include consult note with the request)

Exclusion criteria:

Patients with familial short QT syndrome.

Approval Duration: 3 months

Renewals will be considered on a case-by-case basis

Recommended dose:

200mg administered intravenously or orally every 8 hours for 6 doses followed by a maintenance dose of 200mg daily starting 12 to 24 hours after the last loading dose.

Oral therapy should be considered as a preferred option when clinically appropriate. A loading dose is not required when switching from intravenous to oral treatment or vice versa.

NOTE: Prescribers must submit the laboratory results to confirm the patient's infection and include drugs and drug regimens that have been used for the patient's condition, including the response to prior therapies and other relevant clinical information. Please also confirm that the patient does not meet the exclusion criteria.
For the prophylaxis of cytomegalovirus (CMV) infection in adult patients who have received an allogeneic hematopoietic stem cell transplant (HSCT) meeting the following criteria:

- Age 18 years and older; AND
- Patient is a CMV-seropositive recipient [R+] meeting one of the following circumstances;
  - Recipient using umbilical cord blood as the stem cell source; OR
  - Patient is a haploidentical recipient; OR
  - Recipient of T-cell depleted grafts; OR
  - Recipient with documented history of CMV disease prior to transplantation; OR
  - Recipient requiring high-dose steroids (defined as the use of greater than or the same as 1 mg/kg/day of prednisone or an equivalent dose of another corticosteroid) or other immunosuppression for acute graft versus host disease (GVHD); OR
  - Recipients treated with antithymocyte globulin (ATG) for conditioning, or
  - Recipient treated with ATG for steroid-refractory acute GVHD treatment.
- Patient must have undetectable CMV viremia at baseline (results should be from samples collected within a week of transplant date); AND
- Treatment is prescribed by a clinician with expertise in the management of HSCT (e.g. medical oncologist, hematologist, infectious disease specialist)

Exclusion criteria:
- Treatment of CMV with letermovir is not funded.
- Patients receiving autologous hematopoietic stem cell transplant
- Concomitant use with antiviral drugs used for the management of CMV (e.g. ganciclovir, valganciclovir)

Notes:
- Patients should be transitioned to oral letermovir as soon as clinical circumstances permit to optimize cost-effectiveness

Funded dosage:

A maximum dose of 480mg administered orally or intravenously per day to be started within 28 days of transplant (i.e. as early as the day of transplant and no later than 28 days post-transplant). (240mg when co-administered with cyclosporine)

Approval duration: A maximum duration of funding of 100 days (includes both in-hospital and out-patient utilization) will be provided per patient per HSCT procedure.
For the management of adult cystic fibrosis patients with chronic pulmonary *Pseudomonas aeruginosa* (*P.aeruginosa*) infections who meet the following criteria:

- Documented diagnosis of Cystic Fibrosis; AND
- Patient is 18 years of age or older; AND
- Chronic infection with *Pseudomonas aeruginosa* (PsA) [confirmed by 2 (two) PsA positive sputum cultures taken at least 1 month apart]; AND
- Patient has failed treatment with inhalational tobramycin and demonstrates deteriorating clinical condition despite treatment with inhaled tobramycin; AND
- Request is from a prescriber experienced in the diagnosis and treatment of cystic fibrosis.

Exclusion criteria:

- Use in combination (sequential or cycled during off-treatment periods) with other inhaled antibiotics to treat *P.aeruginosa* will not be funded.
- Funding will not be provided for conditions outside of cystic fibrosis.

Approval duration: 1 year

Renewal requests:

- Patient demonstrates response to therapy.

Approval duration: 1 year
Posaconazole
Brand(s): Posanol
DOSAGE FORM/ STRENGTH: 100 mg tablet, 40 mg/mL Suspension

For the prophylaxis of Aspergillus and Candida infections in patients who have recently (within the past 3 months) undergone an allogeneic bone marrow transplant.

**Duration of Approval:** Limited to 4 months

For the prophylaxis of invasive fungal infections in patients who have previously (3 months or longer) undergone an allogeneic stem cell transplant and are experiencing moderate to severe graft-versus-host-disease (GVHD) will be considered on a case-by-case basis.

Note: Please provide details of the patient's clinical condition including all medications used to treat the condition with your request application.

**Duration of Approval:** Up to 4 months

**Renewals** will be considered on a case-by-case basis for patients who continue to experience ongoing symptoms of moderate to severe GVHD. Please provide information regarding infections that were experienced while on therapy (as applicable) including the names of medications and treatments being used to manage GVHD.

**Duration of Approval:** Case-by-case

For the treatment of invasive aspergillosis* in patients who are refractory or intolerant to voriconazole OR who have documented contraindication to voriconazole.

*Invasive aspergillosis should be confirmed by fungal culture.

Note: Requests without a positive fungal culture must be accompanied by a consultation note from an infectious disease expert with details of how the diagnosis was made and will be considered on a case-by-case basis.

**Duration of Approval:** 3 months

**Renewals** will be considered on a case-by-case basis.
**Posaconazole**

Brand(s): Posanol

**DOSAGE FORM/ STRENGTH:** 100 mg tablet, 40 mg/mL Suspension

For the treatment of mucormycosis** in patients who have failed, have a contraindication to, or experienced intolerance to amphotericin B; OR

**Duration of Approval:** 3 months

For the step-down treatment of mucormycosis** in patients who have been initially treated with amphotericin B but cannot tolerate long-term therapy with this agent.

**Mucormycosis infection must be confirmed by fungal culture.**

*Note: Requests without a positive fungal culture but where the diagnosis of mucormycosis is documented by an infectious diseases consult and other tools (e.g., radiology reports, histopathology, etc.) will be considered on a case-by-case basis.*

**Duration of Approval:** 3 months

**Renewals** will be considered for patients who are responding to therapy but who have not experienced clinical resolution of their condition. Note that requests for renewal must be accompanied by supporting clinical information (Infectious disease consultation/radiology report)

**Duration of Approval:** 3 months

Duration of Approval of subsequent renewal: Case-by-case

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**Valganciclovir**

Brand(s): Valcyte and Generics

**DOSAGE FORM/ STRENGTH:** 450 mg tablet, 50mg/mL pd for oral solution

For the treatment of moderate to severe symptomatic congenital CMV (cCMV) in newborns who meet the following criteria:

- Prescribed **by or in consultation with a pediatric ID specialist** (from one of the 5 treatment centres in Ontario: London, Hamilton, Toronto, Kingston, Ottawa; or Winnipeg for the North Western region of Ontario¹)
- Confirmed diagnosis of cCMV within the first 3 weeks of birth by:
  - PCR (urine, saliva or quantitative serum CMV); OR
  - Positive culture results (urine or saliva)
- Treatment to start within one month of birth
- Evidence of one or more of the following symptoms:
  - CNS disease (e.g., seizures, microcephaly, imaging abnormalities associated with CMV)
  - Eye disease (e.g., chorioretinitis)
  - Severe life-threatening organ dysfunction (specify/describe)
- Regular monitoring of labs for toxicity while on therapy

**Valganciclovir**

*Brand(s): Valtcyte and Generics*

**DOSAGE FORM/ STRENGTH:** 450 mg tablet, 50 mg/mL pd for oral solutions

**Approval Duration:** maximum 6 months at 16mg/kg/dose BID (with dose adjustments in renal dysfunction, < 32 weeks gestational age, etc.)

**Renewals:** No extensions will be provided unless extenuating circumstances for severely affected infants. Case-by-case review with rationale for continued treatment (must include pediatric ID specialist consult note)

All other requests not meeting the above criteria will be reviewed on a case-by-case basis including:
- Initiation of treatment after one month of age
- Evidence of sensorineural hearing loss (SNHL) only (i.e., no other symptom described above)
- Isolated/multiple findings of mild symptoms such as: intrauterine growth retardation (IUGR), thrombocytopenia, elevated liver enzymes, jaundice, hepatitis

¹List of treatment centres and addresses:
- Children's Hospital of Eastern Ontario, 401 Smyth Road, Ottawa ON K1H 8L1
- Kingston General Hospital, 76 Stuart Street, Kingston ON K7L 2V7
- The Hospital for Sick Children, 555 University Avenue, Toronto ON M5G 1X8
- McMaster Children’s Hospital, 1200 Main Street West, Hamilton ON L8N 3Z5
- London Health Sciences Center, 339 Windermere Road, London ON N6A 5A5

**Vancomycin**

*Brand(s): Vancocin and other generics (Note that only specific DINs are reimbursed by the EAP)*

**DOSAGE FORM/ STRENGTH:** 125 mg capsules, 250 mg capsules (on case-by-case basis only)

**Effective September 30, 2019,** Vancomycin oral tablets for the treatment of Uncomplicated Clostridium difficile infection may be accessed upon meeting Limited Use Criteria on the Ontario Drug Benefit Formulary.

Vancomycin Injection to be used as an oral solution for Clostridium Difficile Infection may be accessed through the Telephone Request Service.

Case-by-case consideration for requests not meeting the Limited Use criteria (e.g. higher doses, longer durations, tapering regimens exceeding the dosing limits under LU, complicated C.Difficile Infections) may be considered through external review. Please submit requests to EAP providing adequate and relevant clinical detail to support the request.
**Voriconazole**

Brand(s): VFend  
DOSAGE FORM/ STRENGTH: 50 mg, 200 mg tablets, 200 mg/vial injection

For the treatment of patients who have culture positive candidemia, due to *Candida* species, AND with documented resistance to fluconazole.

This will be for patients whose therapy is initiated in the hospital by a hospital physician and who require continuation of therapy when they are discharged as an outpatient. Oral tablets will be authorized for those with a properly functioning gastrointestinal (GI) tract and the parental injection will be authorized for those who do not have a properly functioning GI.

Case-by-case consideration for other indications will be provided.

**Duration of Approval:** 1 month
ANKYLOSING SPONDYLITIS DRUGS

Adalimumab – See Formulary for funded biosimilars
Brand(s): Humira and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 40mg/0.8mL prefilled syringe, 40mg/0.8mL and 20mg/0.2 mL prefilled pens for subcutaneous injection

Certolizumab
Brand(s): Cimzia
DOSAGE FORM/ STRENGTH: 200 mg/mL prefilled syringe and autoinjector

Etanercept – See Formulary for funded biosimilars
Brand(s): Enbrel and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 25mg/vial and 50mg prefilled syringe for subcutaneous injection

Golimumab
Brand(s): Simponi
DOSAGE FORM/ STRENGTH: 50 mg/0.5 ml prefilled syringe and autoinjector

Infliximab- See Formulary for funded biosimilars
Brand(s): Remicade and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 100mg/10mL intravenous infusion

Secukinumab
Brand(s): Cosentyx
DOSAGE FORM/ STRENGTH: 150 mg/mL prefilled syringe and 150 mg/mL prefilled pen

Reference biologics (e.g. Remicade, Enbrel, Humira) with a provincially funded biosimilar are only considered for provincial funding in patients who are treatment experienced and stable on the reference biologic or those with existing EAP approvals. Prescribers should refer to the ODB formulary for biosimilars and their funded conditions.

It should be noted that after the date when a biosimilar becomes publicly funded for an approved indication, patients initiated on a reference biologic for this same provincially funded indication through support from a manufacturer’s patient support program, may be expected to be provided ongoing access of the reference biologic through the patient support program or to use a biosimilar upon meeting specified criteria. The Ministry will only consider funding of Reference biologics in those who are treatment experienced and stabilized on the product prior to transitioning to the ODB program or in patients with an existing EAP approval.
Refer to the Executive Officer Communications on the Ministry website for Frequently asked questions and notifications of funded biosimilars at http://www.health.gov.on.ca/en/pro/programs/drugs/opdp_eo/eo_communiq.aspx

For the treatment of ankylosing spondylitis (AS) OR psoriatic spondylitis (PS) in patients who have severe active disease with:

- Age of disease onset 50 years of age or younger; AND
- Low back pain and stiffness for greater than 3 months that improves with exercise and not relieved by rest; AND
- Failure to respond to or documented intolerance to adequate trials of 2 non-steroidal anti-inflammatory drugs (NSAIDs) for at least 4 weeks each; AND
- BASDAI score of ≥ 4 for at least 4 weeks while on standard therapy; AND
- A list of current concomitant medications related to the AS/PS, including pain medications (if relevant) with dosing regimens provided.

*NSAIDs include coxibs; use of DMARDS instead of NSAIDs not acceptable

The information submitted with the request must include the following:

- A list of current concomitant medications related to the AS/PS, including pain medications (if relevant). Please include dosing regimens.
- Details of review of radiographic reports for severe active disease.
  - X-ray or CT scan report stating the presence of “SI joint fusion” or “SI joint erosion” OR
  - MRI report stating the presence of “inflammation” or “edema” of the SI joint
  - Actual radiographic reports must be submitted with the request. If the radiographic reports do not specify the above, the request will be reviewed by external medical experts.

Additional information that should be provided if applicable:

- Schober measurement and chest expansion measurement
- Evidence of restricted spinal mobility
- If the patient has AS/PS with predominantly peripheral joint involvement, additional information pertaining to trials of DMARDs must be provided, and these requests will be reviewed by external medical experts.

Duration of Approval: 1 year
Renewal will be considered for patients with objective evidence of at least a 50% reduction in BASDAI score or ≥ 2 absolute point reduction in BASDAI score. Please provide an update on concomitant medications for AS/PS and whether there has been a reduction in pain medication for AS/PS since initiating the biologic (if applicable).

For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided.

The planned dosing regimen for the requested biologic should be provided.

The recommended doses for the treatment of AS/PS are:

- Adalimumab 40 mg every two weeks
- Certolizumab 400mg at 0, 2, and 4 weeks followed by maintenance therapy of 200 mg every 2 weeks or 400 mg every 4 weeks.
- Etanercept 25 mg twice weekly or 50 mg once weekly
- Golimumab 50mg once a month
- Infliximab 3-5mg/kg/dose at 0, 2 and 6 weeks followed by maintenance therapy of up to 5mg/kg/dose every 6 to 8 weeks
- Secukinumab 150 mg sc at weeks 0, 1, 2, and 3 followed by monthly maintenance dosing starting at week 4.

Duration of Approval: First renewal: 1 year, Second and subsequent renewals: 5 years
ASTHMA

Zafirlukast
Brand(s): Accolate
DOSAGE FORM/ STRENGTH: 20 mg tablet

Montelukast
Brand(s): Singulair
DOSAGE FORM/ STRENGTH: 5 mg, 10 mg tablet

For the treatment of asthma patients who cannot manage the use of an inhalation device despite assistance with a spacer (e.g. physically or mentally disabled patients or pediatric patients).

Duration of Approval: 5 years

OR

For the treatment of asthma in children and adolescents whose asthma cannot be controlled on ICS alone and where the condition remains uncontrolled despite using full doses of ICS with addition of LABA, and with assurance of good adherence and inhaler technique

Duration of Approval: 5 years (up until age of 18)

Renewal of requests that meet the above criteria will be provided where the following apply:

• Current medications and dosages must be clearly specified; AND

Objective evidence of positive response from treatment (spirometry OR decrease in health care utilization) must be provided

Duration of Approval: 5 years (up until age of 18)
For the treatment of severe uncontrolled asthma in patients who meet the following criteria:

- Has within the past 12 months required hospitalization for asthma OR has required two or more urgent visits for asthma to a physician or an emergency department OR has had two or more courses of high-dose oral corticosteroids in the past 12 months; AND
- Is age 12 years or older; AND
- Has demonstrated a positive skin test or in vitro reactivity to a perennial aeroallergen; AND
- Has a baseline IgE level between 30 and 700 IU/mL (inclusive); AND
- Has an actual body weight between 20 kg to 150 kg (inclusive); AND
- Is receiving treatment with a high-dose inhaled corticosteroid* in addition to a long-acting inhaled beta 2-agonist. (Note: the patient can be on other concomitant therapies as well); AND
- Is deemed to be adherent and is using his/her inhaled corticosteroid and long-acting beta agonist daily as prescribed; AND
- Is using proper inhaler technique (with a spacer if required); AND
- The request for Xolair is made by the patient’s specialist in respirology or allergy/clinical immunology. (Note: Individual consideration can be given for extenuating circumstances where access to these specialists is not possible.)

* High-dose inhaled corticosteroids are considered the use of more than 1000 mcg of beclomethasone dipropionate (BDP) equivalents daily.
Omalizumab
Brand(s): Xolair
DOSAGE FORM/ STRENGTH: 150 mg/ vial

To avoid delays in the assessment of the request, physicians should provide the following information within their request submission.

1. The number of hospitalizations for asthma in the past 12 months.

2. The number of asthma exacerbations requiring urgent visits to a physician or emergency department in the past 12 months.

3. The average number of night-time awakenings in a one week period. (reflective of control in last 12 months).

4. The average number of puffs/day of short-acting beta-agonists within a one week period (reflective of control in last 12 months).

5. The number of courses of prednisone (or acute increases in prednisone dose if the patient is already using chronic daily prednisone) for asthma exacerbation in the past 12 months.

6. The FEV₁ pre and post bronchodilator.


8. The serum IgE level.

9. Results of a positive allergy testing by skin prick test or IgE RAST.

10. A list of all of the patient’s current asthma medications including drug name and doses.

11. Confirmation that the patient’s asthma is currently uncontrolled despite optimal therapy (including confirmation of proper inhaler technique), patient adherence to current therapy, and the removal of allergic and environmental triggers or the reduction of such triggers to the fullest extent possible.
Omalizumab
Brand(s): Xolair
DOSAGE FORM/ STRENGTH: 150 mg/ vial

Note that contraindications and intolerance to inhaled corticosteroids and/or long-acting beta agonists will not be considered as a justification to request Xolair funding.

Duration of Approval: 1 year

Renewal of requests for Xolair will be considered in patients who have a positive clinical response to the drug and who are expected to continue to do so. Renewals will be considered on a case-by-case basis and should be accompanied by the following information:

1. The number of hospitalizations for asthma in the past 12 months
2. The number of asthma exacerbations requiring urgent visits to a physician or Emergency Department in the past 12 months
3. The number of courses of prednisone (or acute increases in prednisone dose if patient is already using chronic daily prednisone) for asthma exacerbations in the past 12 months.
4. The number of nighttime awakenings (over a several week period post-introduction of therapy)
5. The average number of puffs/day of short-acting beta-agonists used per day (over a several week period post-introduction of therapy)
6. The FEV₁ pre and post bronchodilator
7. All current asthma medications taken by the patient including drug names and dosing schedule.

Duration of Approval: Up to 1 year
Benralizumab
Brand(s): Fasenra
DOSAGE FORM/ STRENGTH: 30 mg/mL Injection (PFS), 30 mg/ml Autoinjector
Updated: April 12, 2021

For the treatment of severe eosinophilic asthma in adult patients who meet ALL the following criteria:

- a) Patient is at least 18 years old; AND
- b) Benralizumab is being used as an add-on maintenance therapy; AND
- c) Patient is inadequately controlled with high dose inhaled corticosteroids, defined as greater than or equal to 500mcg of fluticasone propionate or equivalent daily, and one or more additional asthma controller(s), such as long-acting beta-agonists; AND
- d) Patient has a blood eosinophil count equal to or greater than 300 cell/µL within the past 12 months, and has experienced two or more clinically significant asthma exacerbations, defined as worsening of asthma resulting in administration of systemic corticosteroids for at least 3 days, or an emergency room visit, or hospitalization, in the past 12 months; OR

Patient has a blood eosinophil count of equal to or greater than 150 cells/µL, and is receiving maintenance treatment with oral corticosteroids, defined as greater than the equivalent of prednisone 5 mg per day; AND

- e) Patient is not using any other biologics to treat asthma, such as mepolizumab or omalizumab; AND
- f) Patient has completed a baseline assessment of asthma symptom control using a validated asthma control questionnaire such as the Asthma Control Questionnaire (ACQ) or the Asthma Control Test (ACT) prior to initiation with Fasenra.
- g) Request is submitted by a specialist in respirology, or allergy/clinical immunology, or by a physician with expertise in the treatment of asthma, unless the request confirms that the Patient does not have access to such specialists, in which case the request may be submitted by an Authorized Prescriber

Approved dose for initiation: 30mg sc injection administered once every 4 weeks for the first 3 doses followed by once every 8 weeks thereafter.

Approval duration of initiation and initials: 1 year

Renewals will be considered on a case-by-case basis for patients who do not meet any of the following stopping criteria:

- Patient’s 12-month follow-up asthma control questionnaire score using the same validated asthma control questionnaire completed at baseline does not show improvement, such as a decrease of less than 0.5 points on the ACQ or an increase of less than 3 points on the ACT;
• Patient’s subsequent asthma control questionnaire scores have not been maintained at the score seen following the first 12-months of therapy;

Benralizumab
Brand(s): Fasenra
DOSAGE FORM/ STRENGTH: 30 mg/mL Injection (Autoinjector and Prefilled syringe)

• the number of clinically significant exacerbations, defined as worsening of asthma resulting in administration of systemic corticosteroids for at least 3 days, an emergency room visit or hospitalization, has increased within the previous 12 months; and
• Patient failed to achieve or maintain at least a 25% decrease from baseline in the maintenance oral corticosteroid dose at 12 months of initiation of benralizumab and thereafter.

Approved dose for renewals: 30mg sc once every 8 weeks.

Approval duration for renewals: 1 year

Mepolizumab
Brand(s): Nucala
DOSAGE FORM/ STRENGTH: 100 mg/mL Injection (Vial, Prefilled syringe, Prefilled Autoinjector)

For the treatment of severe eosinophilic asthma in adult patients who meet ALL the following criteria:

i) Patient is at least 18 years old; AND
ii) Mepolizumab is being used as an add-on maintenance therapy; AND
iii) Patient is inadequately controlled with high dose inhaled corticosteroids, defined as greater than or equal to 500mcg of fluticasone propionate or equivalent daily, and one or more additional asthma controller(s), such as long-acting beta-agonists; AND
iv) Patient has a blood eosinophil count equal to or greater than 300 cell/µL within the past 12 months, and has experienced two or more clinically significant asthma exacerbations, defined as worsening of asthma resulting in administration of systemic corticosteroids for at least 3 days, or an emergency room visit, or hospitalization, in the past 12 months;

OR

Patient has a blood eosinophil count of equal to or greater than 150 cells/µL, and is receiving maintenance treatment with oral corticosteroids, defined as greater than the equivalent of prednisone 5 mg per day; AND
v) Patient is not using any other biologics to treat asthma, such as benralizumab or omalizumab; AND

Mepolizumab  
Brand(s): Nucala  
DOSAGE FORM/ STRENGTH: 100 mg/mL Inj. (Vial, Prefilled syringe, Prefilled Autoinjector)

vi) Patient has completed a baseline assessment of asthma symptom control using a validated asthma control questionnaire such as the Asthma Control Questionnaire (ACQ) or the Asthma Control Test (ACT) prior to initiation with Nucala.

vii) Request is submitted by a specialist in respirology, or allergy/clinical immunology, or by a physician with expertise in the treatment of asthma, unless the request confirms that the Patient does not have access to such specialists, in which case the request may be submitted by an Authorized Prescriber

Initial approval duration: 1 year

Renewals will be considered on a case-by-case basis for patients who do not meet any of the following stopping criteria:

a) Patient’s 12-month follow-up asthma control questionnaire score using the same validated asthma control questionnaire completed at baseline does not show improvement, such as a decrease of less than 0.5 points on the ACQ or an increase of less than 3 points on the ACT;

b) Patient’s subsequent asthma control questionnaire scores have not been maintained at the score seen following the first 12-months of therapy;

c) the number of clinically significant exacerbations, defined as worsening of asthma resulting in administration of systemic corticosteroids for at least 3 days, an emergency room visit or hospitalization, has increased within the previous 12 months; and

d) Patient failed to achieve or maintain at least a 25% decrease from baseline in the maintenance oral corticosteroid dose at 12 months of initiation of mepolizumab and thereafter.

Renewal approval duration: 1 year

Ministry Note: The reimbursement criteria for Nucala below were applicable until July 23, 2019. Patients who were reimbursed upon meeting the below criteria may be assessed for renewal of funding meeting this earlier criteria.
Mepolizumab
Brand(s): Nucala
DOSAGE FORM/ STRENGTH: 100 mg/mL Inj. (Vial, Prefilled syringe, Prefilled Autoinjector)

For the treatment of adult patients with severe eosinophilic asthma who meet ALL the following criteria:

a) Mepolizumab is being used as an add-on maintenance therapy; AND

b) Patient is inadequately controlled with high dose inhaled corticosteroids (ICS) (greater than or equal to 500 mcg of fluticasone propionate (FP) equivalents daily) used with one or more asthma controller(s) (e.g., a long-acting beta-agonist [LABA], leukotriene receptor antagonist [LTRA], theophylline); AND

c) Demonstrates a blood eosinophil count greater than or equal to 150 cell/mcL prior to start of mepolizumab (levels must be drawn within 3 months prior to the start of treatment) OR a count greater than or equal to 300 cell/mcL within the 12 months prior to start of mepolizumab; AND

d) Patient must have experienced two or more clinically significant asthma exacerbations in the past 12 months and show reversibility on pulmonary function tests/laboratory spirometry (i.e. at least 12% and 200 mL in FEV1 or FVC from baseline/pre-bronchodilator values) OR is being treated with daily oral corticosteroids for their asthma; AND

e) Request is from a specialist in respirology, or allergy/clinical immunology, or by a physician with expertise in the treatment of asthma. Individual consideration can be given for extenuating circumstances where access to these specialists is not possible.

Duration of approval: 1 year

Renewals will be considered on a case-by-case basis for patients who do not meet ANY of the stopping criteria.

1Stopping criteria are defined as:

- Failure to achieve a decrease in any clinically significant exacerbations (defined as worsening of asthma resulting in administration of systemic corticosteroids for at least 3 days, or an emergency room visit, or hospitalization) within 12 months of initial funding; OR

- Failure to achieve at least a 25% decrease in the maintenance oral corticosteroid dose at 12 months of initiation of mepolizumab.

Duration of renewals: 1 year

Approved dose: 100 mg subcutaneous every 4 weeks.
BLOOD MODIFIERS

Deferasirox
Brand(s): Exjade
DOSAGE FORM/ STRENGTH: 125 mg, 250 mg, 500 mg tablet
Brand(s): Jadenu
DOSAGE FORM/ STRENGTH: 90 mg, 180 mg, 360 mg tablet

For the treatment of patients with chronic iron overload in transfusion-related anemia due to \( B \)-thalassemia or sickle cell disease in patients who meet the following criteria;

- Patient is 6 years of age or older; OR
- The patient is between 2 to 5 years of age (inclusive) and cannot be adequately treated with deferoxamine.

Combination therapy [i.e. Deferasirox (Jadenu or Exjade) in addition to another iron chelating agent] will not be approved for funding.

Therapy should be initiated and maintained by physicians experienced in the treatment of chronic iron overload due to blood transfusions.

**Renewals** will be considered in patients who continue to require iron chelation therapy and has had a consistent response to therapy (demonstrated by a reduction in baseline LIC levels).

The following documentation is required for renewals:

- A transfusion record from the past year; AND
- LIC levels – baseline (pre-treatment) and since initiation of treatment. The most recent LIC level should be from within the previous year.

Approval duration of initiation: 5 years
Approval duration of renewals: 5 years

For the treatment of chronic iron overload in transfusion-dependent anemia in those with low-risk myelodysplastic syndrome (MDS) or other rare anemias (e.g. Diamond Blackfan) in patients who have a contraindication or severe intolerance to deferoxamine.

Contraindications may include one or more of the following:

- known or suspected hypersensitivity to deferoxamine
- recurrent injection or infusion-site reactions (e.g., cellulitis)
- concomitant bleeding disorder
- immunocompromised patients with a documented risk of significant infections with parenteral administration (e.g. neutropenia)

**Duration of Approval:** 1 year

**Renewals** will be considered on a case-by-case basis. Physicians must provide adequate information to support the request for renewal.
<table>
<thead>
<tr>
<th>Deferasirox</th>
<th>Brand(s): Exjade</th>
<th>DOSAGE FORM/ STRENGTH: 125 mg, 250 mg, 500 mg tablet</th>
</tr>
</thead>
<tbody>
<tr>
<td>Brand(s): Jadenu</td>
<td>DOSAGE FORM/ STRENGTH: 90 mg, 180 mg, 360 mg tablet</td>
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Note: If switching between brands of deferasirox, please include the reasons for switching (e.g. description of the intolerance etc.) with your request application.

**Duration of Approval:** 5 years
Eculizumab
Brand(s): Soliris
DOSAGE FORM/ STRENGTH: 10 mg/mL (300 mg per vial)

For the treatment of patients with Paroxysmal Nocturnal Hemoglobinuria (PNH) meeting the following criteria:

The diagnosis of Paroxysmal Nocturnal Hemoglobinuria (PNH) has been made based on the following confirmatory results:

- Flow cytometry/FLAER exam with granulocytes clone ≥ 10% AND
- LDH > 1.5 ULN

AND at least one of the following:

- A thrombotic or embolic event which required the institution of therapeutic anticoagulant therapy,
- Minimum transfusion requirement of 4 units of red blood cells in the previous 12 months,
- Chronic or recurrent anemia where causes other than hemolysis have been excluded and demonstrated by more than one measure of less than or equal to 70 g/L or by more than one measure of less than or equal to 100 g/L with concurrent symptoms of anemia,
- Pulmonary insufficiency: Debilitating shortness of breath and/or chest pain resulting in limitation of normal activity (New York Heart Association Class III) and/or established diagnosis of pulmonary arterial hypertension, where causes other than PNH have been excluded,
- Renal insufficiency: History of renal insufficiency, demonstrated by an eGFR less than or equal to 60 mL/min/1.73 m², where causes other than PNH have been excluded,
- Smooth muscle spasm: Recurrent episodes of severe pain requiring hospitalization and/or narcotic analgesia, where causes other than PNH have been excluded.

The dose of eculizumab that will be considered is:

600 mg once per week for the first 4 weeks, then from week five of treatment, 900 mg once every 2 weeks

Duration of Approval: 6 months
Eculizumab
Brand(s): Soliris
DOSAGE FORM/ STRENGTH: 10 mg/mL (300 mg per vial)

Renewals will be considered for patients who;

- Demonstrate clinical improvement while on therapy or
- Where therapy has been shown to stabilize the patient’s condition

Requests for renewal should be accompanied by confirmation of granulocyte clone size (by flow cytometry).

Further, subsidized treatment may continue unless one or more of the following situations apply:

i) The patient or treating physician fails to comply adequately with treatment or measures, including monitoring requirements, taken to evaluate the effectiveness of the therapy;

ii) If therapy fails to relieve the symptoms of disease that originally resulted in the patient being approved for subsidized treatment;

Other eligibility requirements:

Note: All patients must receive meningococcal vaccination with a tetravalent vaccine at least two weeks prior to receiving the first dose of eculizumab.

Exclusion criteria for both initial and renewal requests:

i) Small granulocyte clone size - the treatment of patients with a granulocyte clone size below 10% will not be eligible for treatment; OR

ii) Aplastic anemia with two or more of the following: neutrophil count below 0.5 x 10⁹/L, platelet count below 20 x 10⁹/L, reticulocytes below 25 x 10⁹/L, or severe bone marrow hypocellularity; OR

iii) Patients afflicted with PNH and another life-threatening or severe disease where the long term prognosis is unlikely to be influenced by therapy (for example acute myeloid leukemia or high-risk myelodysplastic syndrome); OR

iv) The presence of another medical condition that might reasonably be expected to compromise a response to therapy.
Eculizumab
Brand(s): Soliris
DOSAGE FORM/ STRENGTH: 10 mg/mL (300 mg per vial)

Preamble:
A confirmed diagnosis of atypical hemolytic uremic syndrome (aHUS) is required for eculizumab funding. The information below is to provide clinicians with context for how a diagnosis of aHUS will be assessed for funding consideration. Details to address these issues should be provided in the funding request.

While some patients may already have a confirmed aHUS diagnosis, by clinical history and/or genetic testing, the majority of patients presenting with thrombotic microangiopathy (TMA) have no prior diagnosis of aHUS. For most patients presenting with a TMA, it is not possible to confidently separate aHUS from the vast majority of other conditions causing TMA until after appropriate testing and treatment have occurred. The majority of patients who have TMA suffer from Thrombotic Thrombocytopenic Purpura (TTP) (30-40%), or a secondary form of TMA (e.g., pregnancy, HIV, collagen vascular disease, drugs, malignancy, stem cell transplant, malignant hypertension) (> 50%), or hemolytic uremic syndrome due to a Shiga toxin (>5%). In most cases, patients who suffer from TTP will have an ADAMTS-13 of less than 10%. If TTP has been ruled out and any secondary causes have been treated and the patient still has a persisting unexplained TMA with ADAMTS-13 ≥ 10%, the patient would be presumed to suffer from aHUS. Patients who present with ADAMTS-13 of ≥10% and who are unresponsive to plasma therapy (>4 plasma exchanges) and do not have a known secondary explanation would also be presumed to suffer from aHUS.

In the absence of a confirmed diagnosis of aHUS, there is nothing in these criteria that changes the clinical expectation for appropriate use of plasma exchange/plasma infusion in the management of patients presenting with TMA.

**Initiation Criteria**
A patient must meet all three of the following criteria to obtain funding for initial treatment with eculizumab:

1. Confirmed diagnosis* of atypical hemolytic uremic syndrome (aHUS) at initial presentation, defined by:
   
   a. Presence of an unexplained non-disseminated intravascular coagulation thrombotic microangiopathy (TMA); AND
   
   b. Baseline ADAMTS-13 activity ≥ 10% on blood samples taken prior to plasma exchange or plasma infusion (PE/PI);
Eculizumab
Brand(s): Soliris
DOSAGE FORM/ STRENGTH: 10 mg/mL (300 mg per vial)

Note:

If the sample for ADAMTS-13 was not collected prior to PE or PI, platelet counts > 30 x 10^9/L and eGFR < 50 mL/min/1.73m^2 at TMA presentation will be accepted as predictive of ADAMTS-13 ≥10% in TMA patients. In this case, measurement of ADAMTS-13 can be taken 1-2 weeks following the last PE. The ADAMTS-13 result must be provided within 30 days of commencement of eculizumab and at least 1 week after the last PE. A one-month interim funding for eculizumab will be provided.

AND

c. STEC-negative test in patients with a history of bloody diarrhea in the preceding two weeks; AND
d. Other diagnoses and causes of TMA must be ruled out, as per preamble.

2. Evidence of ongoing active and progressing TMA as defined by:

a. Thrombocytopenia (platelet count <150 x 10^9/L) that is not explained by some other cause including secondary TMA; AND hemolysis as indicated by the documentation of two of the following: red blood cell (RBC) fragmentation (schistocytes) on the blood film; low or absent haptoglobin; or lactate dehydrogenase (LDH) above normal; OR

b. Tissue biopsy confirming TMA in patients who do not have evidence of platelet consumption and hemolysis.

Note: Review by external clinical expert may be required to assess requests for patients with ongoing TMA that may not clearly meet the above criteria.

3. Evidence of at least one of the following documented clinical features of active organ damage or impairment:

a. Kidney impairment as demonstrated by one of the following:
   o A decline in estimated glomerular filtration rate (eGFR) or a rise in serum creatinine (SrCr) of >20% in a patient with pre-existing renal impairment; OR
   o SrCr > upper limit of normal (ULN) for age or eGFR < 60mL/min in patients who have no history of pre-existing renal impairment (i.e., who have no baseline eGFR measurement); OR
   o SrCr > the age-appropriate ULN in pediatric patients (subject to advice from a pediatric nephrologist); or
**Eculizumab**
Brand(s): Soliris
**DOSAGE FORM/ STRENGTH:** 10 mg/mL (300 mg per vial)

- Renal biopsy;

OR

b. Onset of neurological impairment related to TMA (e.g., visual field defect, hemiparesis, sensory loss, asymmetric limb weakness, confusion, loss of consciousness/coma, new onset seizure).

Note: Patients who have extra-renal complications related to TMA (e.g., TMA-related cardiac impairment, TMA-related gastrointestinal impairment, or TMA-related pulmonary impairment) will be reviewed by an external clinical expert.

**Continuation Criteria (at 6 months)**

After six months of eculizumab therapy, a further six month of funding will be considered if the patient demonstrates treatment response, defined as:

Hematological normalization (platelet count, LDH, haptoglobin); AND

- An improvement or stabilization of eGFR (or SrCr); AND
- Stabilization of neurological or extra-renal impairment if these complications were originally present.

Continued treatment with eculizumab will not be funded beyond six months if a patient has experienced treatment failure, defined as:

- Dialysis-dependent at six months, and failed to demonstrate resolution or stabilization of neurological or extra-renal complications if these were originally present; OR
- On dialysis for ≥ four of the previous six months while receiving eculizumab and failed to demonstrate resolution or stabilization of neurological or extra-renal complications if these were originally present; OR
- Worsening of kidney function with a reduction in eGFR or increase in SrCr ≥ 25% from baseline.

Approval duration: 6 months

**Continuation Criteria (at 12 months):**

1. Ongoing treatment response as defined in the 6-month continuation criteria; AND
2. The patient has limited organ reserve defined as:
Eculizumab
Brand(s): Soliris
DOSAGE FORM/ STRENGTH: 10 mg/mL (300 mg per vial)

- Significant cardiomyopathy, neurological, gastrointestinal or pulmonary impairment related to TMA; or
- Grade 4 or 5 chronic kidney disease (eGFR <30mL/min). (Note: Patients who are dialysis-dependent with no significant extra-renal manifestations persisting are not considered).

There may be other exceptional circumstances where the patient has a high risk of recurrence and in whom consequences of a relapse are significant (e.g., complement Factor H genetic mutation, multiple clinical presentations of active TMA). These will be reviewed on a case-by-case basis by an external clinical expert.

For patients in whom a pause in therapy is recommended, funding will be left in place for 3 months so that eculizumab can be quickly restarted upon evidence of recurrence per recommencement criteria.

Approval duration: 12 months

Recommencement Criteria:

A patient previously diagnosed with aHUS and who responded to treatment with eculizumab and has not failed eculizumab is eligible to restart eculizumab if the following clinical conditions are met:

- Significant hemolysis as evidenced by presence of schistocytes on the blood film, or low or absent haptoglobin, or LDH above normal;

AND EITHER

- Platelet consumption as measured by either ≥ 25% decline from patient baseline or thrombocytopenia (platelet count <150,000 x 10^9/L);

OR

- TMA-related organ impairment (e.g., unexplained rise in serum creatinine with onset of urine dipstick positive for hemoglobin) including on recent biopsy.

Note:

1. Raised LDH alone is not a sufficient reason to recommence eculizumab, but thrombocytopenia with one marker of hemolysis (such as raised LDH, presence of schistocytes, or low/absence of haptoglobin) is an accepted reason to recommence.

2. Kidney transplantation/dialysis is not a contraindication to recommencement.
Eculizumab
Brand(s): Soliris
DOSAGE FORM/ STRENGTH: 10 mg/mL (300 mg per vial)

A patient who becomes eligible to restart eculizumab, in accordance with the above criteria, will be assessed every 6 months for treatment response or failure.

Approval duration: 6 months

Patients undergoing kidney transplantation:

For patients with a confirmed aHUS diagnosis who are undergoing kidney transplantation, eculizumab funding will be provided for the time period immediately prior to (or at time of) transplant. Treatment must be started immediately prior to or at time of transplant.

Approval duration: 6 months

All funding requests must come from, or be submitted in consultation with, a pediatric nephrologist, a nephrologist, a pediatric hematologist or a hematologist.

Deferiprone
Brand(s): Ferriprox
DOSAGE FORM/ STRENGTH: 1000 mg Tablets, 100 mg/mL oral solution

For the treatment of patients with transfusional iron overload due to thalassemia syndromes who cannot be adequately treated with deferoxamine or deferasirox.

Notes:

Combination iron chelation therapy with Ferriprox will be considered on a case-by-case basis.

Therapy should be initiated and maintained by physicians experienced in the treatment of chronic iron overload due to blood transfusions.

Duration of Approval: 5 years

Renewals will be considered for Patients who continue to require iron chelation therapy and has had a consistent response to therapy (demonstrated by a reduction in baseline liver iron concentration (LIC) levels).

The following documentation is required:

- A transfusion record from the past year; and
- LIC levels – baseline (pre-treatment) and since initiation of treatment. The most recent LIC level should be from within the previous year.

Duration of Approval: 5 years
For the treatment of refractory chronic idiopathic thrombocytopenic purpura (ITP) with bleeding complications in patients who meet the following criteria:

- Patient has undergone a splenectomy\(^1\); AND
- Patient has tried and is unresponsive to other treatment modalities\(^2\).

\(^1\)Requests for Revolade where the requesting physician has stated that the patient is not a candidate for splenectomy will be assessed on a case-by-case basis. The requesting physician must provide rationale for why a splenectomy cannot be considered, and where possible, to include a preoperative/surgical evaluation on the patient’s surgical risks to splenectomy, to include consideration of risks of laparoscopic and open surgical interventions if these are available. This evaluation must come from a physician who is not the requesting physician.

\(^2\)Appropriate first-line treatment modalities may include:

- Corticosteroids
- IV anti-D
- Intravenous immune globulin (IVIG)

Appropriate second-line treatment modalities include:

- Azathioprine
- Cyclosporine
- Cyclophosphamide
- Mycophenolate
- Rituximab
- Danazol
- Dapsone
Eltrombopag
Brand(s): Revolade
DOSAGE FORM/ STRENGTH: 25 mg, 50 mg tablet

Note: Patients need to have failed at least two of the second-line therapies listed above prior to requesting Revolade. Dosage: 50 mg once daily to a maximum of 75 mg once daily.

Duration of Approval: 1 year

Renewal of requests for Revolade will be assessed on a case-by-case basis.

Note: Revolade therapy beyond 1 year of continuous treatment has not been studied. After 1 year of continuous treatment, therapeutic options should be reassessed.

Duration of Approval: 1 year
**Icatibant**
Brand(s): Firazyr  
DOSAGE FORM/ STRENGTH: 30 mg/3 mL prefilled syringe

For the treatment of acute attacks of type I or type II hereditary angioedema (HAE) in adults with lab confirmed c1-esterase inhibitor deficiency if the following conditions are met:

- a. Treatment of acute non-laryngeal attacks of at least moderate severity; OR
- b. Treatment of acute laryngeal attacks; AND
- c. Must be prescribed by physicians (e.g. immunologists, allergists or hematologists) with experience in the treatment of HAE.

Notes:

- Documentation of diagnosis (e.g. patient and family history, symptoms, lab test results) must be provided.
- For acute non-laryngeal attacks, documentation of severity (frequency, location, and degree of swelling) must be provided

Doses for acute treatment are limited to a single dose for self-administration per attack.

**Duration of Approval:** Lifetime

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**Lanadelumab**
Brand(s): Takhzyro  
DOSAGE FORM/ STRENGTH: 300 mg/2mL  
Effective date: February 2, 2021

**Initiation Criteria:**

For the routine prevention of attacks of hereditary angioedema (HAE) in patients meeting ALL the following criteria:

- Is at least 12 years of age; AND
- Has HAE type I or II; AND
- Has experienced at least three HAE attacks within any four-week period requiring treatment with an acute injectable therapy (e.g. Icatibant, C1-INH); AND
- Must be prescribed by physicians (e.g. immunologists, allergists or hematologists) with experience in the diagnosis and management of HAE.
**Lanadelumab**
Brand(s): Takhzyro
DOSAGE FORM/ STRENGTH: 300 mg/2mL

NOTE: For patients who are already on long-term prophylactic treatment of angioedema (e.g. C1-INH) and intend to transition to lanadelumab, please provide historical HAE attack rates and treatment details.

**Discontinuation Criteria:**

Reimbursement will be discontinued in patients who either respond inadequately or exhibit loss of response, defined as follows:

- **Inadequate response:** No reduction in the number of HAE attacks that require treatment with an acute injectable therapy during the first three months of treatment with Takhzyro.
- **Loss of response:** An increase in the observed number of HAE attacks requiring acute injectable treatment compared to the number before initiating treatment with lanadelumab.

**Exclusion Criteria:** Patient is using lanadelumab with other medications used for the long-term prophylactic treatment of angioedema (e.g. C1-INH).

**Renewal Criteria:** Renewals will be considered in patients who do not meet the discontinuation criteria.

**NOTE(S):**

- *An assessment of a response to treatment should be conducted three months after initiating treatment with lanadelumab.*
- *Following the initial three-month assessment, patients should be assessed for continued response to lanadelumab every six months.*

**Recommended dose:** 300 mg administered subcutaneously (SC) every 2 weeks. The dose should not be escalated to more than 300 mg every two weeks in cases of inadequate response or loss of response. Dosing interval of 300 mg SC every 4 weeks may be considered if the patient is well-controlled (e.g. attack free) for more than six months.

**Approval duration of initials and renewals:** 4 months initial, 6 months first renewal, 12 months subsequent renewals

**Case by Case Consideration:** for patients not meeting the requirement of 3 HAE attacks in the initiation criteria but with HAE attacks associated with significant functional impairment and risk of mortality (e.g. laryngeal attacks).
Romiplostim
Brand(s): Nplate
DOSAGE FORM/ STRENGTH: 250 mcg/0.5 mL 500 mcg/mL

For the treatment of refractory chronic idiopathic thrombocytopenic purpura (ITP) with bleeding complications in patients who meet the following criteria;

i) Patient has undergone a splenectomy¹

ii) Patient has tried and is unresponsive to other treatment modalities².

¹Requests for romiplostin where the requesting physician has stated that the patient is not a candidate for splenectomy will be assessed on a case-by-case basis. The requesting physician must provide rationale for why a splenectomy cannot be considered, and where possible, to include a preoperative evaluation on the patient’s surgical risks to splenectomy to include consideration of risks of laparoscopic and open surgical interventions if these are available.

Note: The Executive Officer (EO) may revise the criteria if the frequency of patients who are not eligible for splenectomy exceeds published estimates.

²Appropriate first-line treatment modalities may include;
- Corticosteroids
- IV anti-D
- Intravenous immune globulin (IVIG)

²Appropriate second-line treatment modalities may include;
- Azathioprine
- Cyclosporine
- Cyclophosphamide
- Mycophenolate
- Rituximab
- Danazol
- Dapsone

Patients need to have failed at least two second-line therapies prior to requesting Nplate.

Duration of Approval: 1 year

Renewal of requests will be considered in patients who have a stable platelet response and reduced symptoms of ITP-related bleeding events.

Duration of Approval: 1 year
Eplerenone
Brand(s): Inspra
DOSAGE FORM/ STRENGTH: 25 mg, 50 mg tablets

For the treatment of patients who have heart failure and left ventricular systolic dysfunction due to acute myocardial infarction. Patients must have:

- An ejection fraction ≤ 40% AND
- Prior trial of spironolactone but experienced severe symptomatic (painful) gynecomastia

Duration of Approval: Lifetime
Edaravone
Brand(s): Radicava
DOSAGE FORM/ STRENGTH: 30mg/100mL bags

Initiation Criteria:
For the treatment of amyotrophic lateral sclerosis (ALS) in patients meeting ALL the following criteria:

- Diagnosis of definite ALS or probable ALS; AND
- ALS symptom onset occurred within the past two years or less; AND
- Has a score greater than or the same as two (2) points on each of the 12 items of the ALS Functional Rating Scale – Revised (ALSFRS-R); AND
- Forced vital capacity (FVC) is greater than or equal to 80% of predicted; AND
- Does not require permanent non-invasive or invasive ventilation; AND
- Is under the care of a specialist with experience in the diagnosis and management of ALS

Discontinuation Criteria:
Reimbursement will be discontinued in patients who meet any one of the following criteria:

- patient becomes non-ambulatory (ALSFRS-R score ≤ 1 for item 8) AND is unable to cut food and feed themselves without assistance, irrespective of whether a gastrostomy is in place (ALSFRS-R score < 1 for item 5a or 5b); or
- patient requires permanent non-invasive or invasive ventilation.

Renewal Criteria:
Renewals will be considered in patients who do not meet the discontinuation criteria.

Recommended dose:
60 mg administered as an intravenous infusion according to the following schedule:

- An initial treatment cycle with daily doses for 14 days, followed by a 14-day drug-free period
- Subsequent treatment cycles with daily doses for 10 days out of 14-day periods, followed by 14-day drug-free periods.

Approval duration of initials and renewals: 1 year
**Modafinil**  
Brand(s): Alertec  
**DOSAGE FORM/ STRENGTH:** 100 mg tablet

For the symptomatic treatment of excessive daytime sleepiness in patients with **narcolepsy** who have demonstrated a lack of response to or an inability to tolerate dextroamphetamine AND methylphenidate.

Note: See also Multiple Sclerosis Drugs

**Duration of Approval:** 2 years (Initials and Renewals)

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**Riluzole**  
Brand(s): Rilutek  
**DOSAGE FORM/ STRENGTH:** 50 mg tablet

Approvals will be provided for:

Patients who have probable or definite amyotrophic lateral sclerosis (ALS) as defined by World Federation of Neurology (WFN) criteria with onset within 5 years, who have a vital capacity of >60% predicted and do not have a tracheostomy.

**Discontinuation Criteria:**

Reimbursement will be discontinued if the patient progresses to require permanent assisted ventilation. This is defined as assisted ventilation required for 23 out of 24 hours for greater than or equal to 14 consecutive days.

**Renewal Criteria:**

Renewals will be considered in patients who do not meet the discontinuation criteria.

**Approval period of initials and renewals:** 12 months

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**Tetrabenazine**  
Brand(s): Nitoman  
**DOSAGE FORM/ STRENGTH:** 25 mg tablet

For the treatment of Huntington’s chorea, tic and Gille’s de la Toureet syndrome and tardive dyskinesia in patients meeting the following criteria:

i) is prescribed by (or in consultation with) physicians who are experienced in the treatment of hyperkinetic movement disorders (e.g. specialists practicing in a Movement Disorder Clinic, neurologists, psychiatrists, physiatrists, geriatricians, pediatricians); **AND**
ii) have disabling Huntington’s chorea OR tic and Gille’s de la Tourette syndrome and have documented evidence of failure to respond, intolerable side effects or contraindication to at least one agent presently available on the Formulary.

**Tetrabenazine**
Brand(s): Nitoman
DOSAGE FORM/ STRENGTH: 25 mg tablet

**Note that for patients with disabling tardive dyskinesia, a trial of a Formulary agent is NOT required (ie. tetrabenazine can be considered for use as a first-line agent)**

Duration of Approval: 1 year

**Renewals** will be considered for patients whose request is prescribed by (or in consultation with) physicians who are experienced in the treatment of hyperkinetic movement disorders (e.g. specialists practicing in a Movement Disorder Clinic, neurologists, psychiatrists, physiatrists, geriatricians, pediatricians); AND who provide written confirmation that movements and functional status are stabilized on tetrabenazine therapy.

Duration of Approval: 5 years

For the treatment of Hemiballismus, senile chorea, or other disabling hyperkinetic movement disorders (HKMD) will be considered on a case-by-case basis in patients meeting the following criteria:

- is prescribed by (or in consultation with) physicians who are experienced in the treatment of hyperkinetic movement disorders (e.g. specialists practicing in a Movement Disorder Clinic, neurologists, psychiatrists, physiatrists, geriatricians, pediatricians); **AND**

- have documented evidence of failure to respond, intolerable side effects or contraindication to at least one agent presently available on the Formulary.

Duration of Approval: 1 year

**Renewals** will be considered for patients whose request is prescribed by (or in consultation with) physicians who are experienced in the treatment of hyperkinetic movement disorders (e.g. specialists practicing in a Movement Disorder Clinic, neurologists, psychiatrists, physiatrists, geriatricians, pediatricians); AND who provide written confirmation that movements and functional status are stabilized on tetrabenazine therapy.

Duration of Approval: 5 years

Please note that information MUST BE provided about why a patient has not tried or cannot try a formulary alternative.
Requests not meeting the above criteria for HKMD will be considered through a case-by-case review and the physician must provide adequate clinical information to enable this assessment.
For the treatment of adult patients with active moderate to severe hidradenitis suppurativa who have not responded to conventional therapy (including systemic antibiotics) and who meet all of the following:

- 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
- Prescribed by a practitioner with expertise in the management of patients with HS

Note: Treatment with adalimumab should be discontinued if there is no improvement after 12 weeks of treatment

First renewal:

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

Subsequent renewal:

- For renewals beyond the second year, objective evidence of the preservation of treatment effect should be provided (i.e. the current AN (abscess and inflammatory nodule) count and draining fistula count should be compared to the count prior to initiating treatment with adalimumab).

Approval duration:

- Initial approval: 3 months
- First renewals: 1 year
- Subsequent renewals: 2 years

Recommended dose:

- The recommended dose is 160 mg initially (week 0), followed by 80 mg at week 2, then 40 mg at week 4, and 40 mg weekly thereafter
**Initiation criteria**

For the treatment of moderate to severe atopic dermatitis in patients meeting the following criteria;

1. 12 years of age or older; AND
2. Diagnosed with moderate-to-severe atopic dermatitis (AD) as defined by an Eczema Area and Severity Index (EASI) score equal to or greater than 16 points; AND
3. Failure to maintain adequate control of their dermatitis with topical prescription therapies (if these therapies are not advisable, please provide reasons why they could not be used with your application); AND
4. Failure or intolerance with an adequate trial of phototherapy (where available), methotrexate, and cyclosporine$^1$; AND
5. Dupilumab is prescribed by a dermatologist, pediatrician or immunologist, or in consultation with one of these specialists. (Please include the consult note with the EAP application.)

Notes:

$^1$For each treatment used, provide documentation of refractory disease and/or intolerance (including a description of the adverse effect and severity of reaction). If a patient is deemed to be ineligible or contraindicated to receive the treatment, provide the reason(s) for their ineligibility.

$^2$The concurrent use of dupilumab used in combination with other biologics used for other conditions will be considered on a case-by-case basis.

**Exclusion Criteria:** (Patients meeting any of the following will not be funded)

1. Dupilumab will not be funded if it is being used in combination with phototherapy or immunosuppressant drugs (e.g. methotrexate, cyclosporine)
2. Dupilumab will not be funded in combination with other biologics used for atopic dermatitis$^2$.

**Renewal Criteria:**

**First renewal:**

Renewal of funding will be considered in patients with documentation of benefit from treatment. Benefit from treatment is defined as a 75% or greater improvement from
baseline in the Eczema Area and Severity Index (EASI) score (EASI-75) within six months of treatment initiation.

**Dupilumab**

**Brand(s):** Dupixent  
**DOSAGE FORM/ STRENGTH:** 200mg/1.14mL, 300mg/2mL Injection solution

**Subsequent renewals:**

Subsequent renewal of funding will be considered in patient who maintain the 75% or greater improvement in EASI-75 response from baseline.

**Duration of Approval for initial requests:** 6 months  
**Duration of Approval for first and second renewal:** 6 months  
**Duration of Approval for 3rd and subsequent renewals:** 1 year

**Recommended dose:**

Adults: An initial dose of 600 mg followed by 300 mg every other week.

Refer to the product monograph for dosing in adolescents 17 year of age and younger.

**Imiquimod**

**Brand(s):** Apo-Imiquimod  
**DOSAGE FORM/ STRENGTH:** 5% Cream

**For the treatment of external genital and perianal warts/condyloma acuminata** in patients who;

- Have documented failure to a trial of podophyllum resin and one other treatment modality (including cryotherapy, surgical excision, or electrosurgery).

**Duration of Approvals:** 1 year (Maximum of 16 weeks for each treatment course )

**For the treatment of biopsy-confirmed primary superficial basal cell carcinoma** in patients meeting the following criteria;

- Tumour diameter of ≤ 2 cm AND
- Tumour location on the trunk, neck or extremities (excluding hands and feet) AND
- Surgery or irradiation therapy is not medically indicated (e.g. recurrent lesions in previously irradiated area, number of lesions too numerous to irradiate or remove surgically)
Duration of Approvals: 6 weeks

Renewals for the same tumour will not be considered.
Omalizumab
Brand(s): Xolair
DOSAGE FORM/ STRENGTH: 150 mg Inj

Initial Criteria:

For the treatment of moderate to severe chronic idiopathic urticaria (CIU) when prescribed by a specialist (i.e. an allergist, an immunologist, a dermatologist) in patients who meet ALL the following criteria;

i) Patient must be 12 years of age or older; AND
ii) Patient must remain symptomatic despite optimum management with available oral therapies.

Approved regimen: Up to 300 mg every 4 weeks

Duration of Approval: 24 weeks

Renewals will be considered for patients who demonstrate one of the following responses to treatment;

i) Patient has had a trial of stopping omalizumab treatment after having achieved symptom control for at least 12 weeks while on therapy but who experience symptom relapse during the stoppage period; OR
ii) Patient has demonstrated improvement but has not been able to achieve complete symptom control for more than 12 consecutive weeks; OR
iii) Patient has demonstrated a partial response to treatment defined as at least a greater than or equal to 9.5 point reduction in the baseline urticaria activity score over 7 days (UAS7).

Approved regimen: Up to 300 mg every 4 weeks

Duration of Approval of Renewals: 24 week
Propranolol
Brand(s): Hemangiol
DOSAGE FORM/ STRENGTH: 3.75mg/mL oral solution

Initial Criteria:

For the treatment of infants and children with any of the following proliferating infantile hemangiomas;

- Life or function-threatening hemangioma OR
- Ulcerated hemangioma in those experiencing pain and/or lack of response to simple wound care measures; OR
- Hemangiomas deemed to put the patient at risk of permanent scarring or disfigurement.

Requests must be from a dermatologist or a physician experienced in the care of infantile hemangiomas.

Duration of approval: up to 12 months¹

¹ Note that the treatment duration is typically 6 months and consideration should be made to discontinue the product in the absence of improvement within the first 2 months.

Renewals: Renewals will be considered on a case-by-case basis. If wounds are not healing, please provide clinical information as to why ongoing reimbursement is required.
Rituximab – see Formulary for funded biosimilars

Brand(s): Riximyo, Ruxience, Truxima (Biosimilars); Rituxan (Originator)

DOSAGE FORM/ STRENGTH: 10 mg/mL intravenous injection

For the treatment of severe pemphigus vulgaris in patients who meet the following criteria.

- Patient has failed combination therapy with high-dose systemic steroids\(^1\) and a steroid-sparing immunosuppressant\(^2\) trialed in combination for a minimum of 3 months.

- The request must be made by a dermatologist/specialist familiar with the management of pemphigus vulgaris and with the use of rituximab in this condition.

\(^1\)Patients must have used a steroid dose equivalent to a 1 mg/kg prednisone dose equivalent (or a minimum of 60 mg/day for patients > 60 kg) for at least 4 to 6 weeks before attempting to taper to a lower dose.

\(^2\)Patients must try at least one of the following at therapeutic doses: azathioprine, mycophenolate, cyclophosphamide, or methotrexate (in combination with a steroid).

Dose: ONE course of treatment with rituximab is considered

375 mg/m\(^2\) administered weekly for 4 weeks (for a total of 4 doses) OR

1000 mg of rituximab administered at week 0 and week 2 (for a total of 2 doses)

Re-treatment may be provided if the patient responded to rituximab therapy then experiences disease flare, as long as the request is made no less than 6 months after the last dose of the patient’s last treatment course/cycle with rituximab.

Rejection Criteria:

- Other dermatology diagnoses, such as pemphigus foliaceus and bullous pemphigoid

- Maintenance infusions (i.e. regular maintenance doses to keep disease in remission)

Duration of Approval: 1 year

Maintenance Treatment is not funded.

First Renewal: 1 year

Subsequent Renewals after first renewal: 2 years

(Rituximab is funded for course of therapy to be given at an interval of at least 6 months only upon flare of the condition.)
ENDOCRINOLOGY AGENTS

**Cabergoline**
Brand(s): Dostinex and generics (see formulary for OFI listings)
DOSAGE FORM/ STRENGTH: 0.5 mg tablets

For the treatment of hyperprolactinemia.

Note that cabergoline may be used as any line of therapy for hyperprolactinemia (i.e. first or subsequent lines)

Approval duration: Lifetime

**Cinacalcet**
Brand(s): Sensipar
DOSAGE FORM/ STRENGTH: 30 mg, 60 mg, 90 mg tablets

For the treatment of severe hyperparathyroidism* in patients with chronic kidney disease who are on dialysis who meet the following criteria;

i) the patient is refractory to other treatments; AND

ii) the patient has symptoms clearly related to hyperparathyroidism that are causing significant impairment in quality of life (e.g. calciphylaxis or bone pain); AND

iii) additionally, ONE of the following criteria is present:

- the patient has been reviewed by a surgeon, anesthetist or nephrologist and has been deemed to not be a candidate for parathyroidectomy due to high surgical risk or anesthetic risk. [Please note: This must be accompanied by a clinical note explaining the high surgical risk or anesthetic risk and the patient’s parathyroid hormone (PTH) level]; OR

- the patient has been wait-listed for a parathyroidectomy and requires Sensipar for bridge therapy; OR

- the patient is awaiting an imminent renal transplant and a nephrologist indicates a preference for pre-transplant treatment with Sensipar instead of a parathyroidectomy.

*Severe hyperparathyroidism is considered to be patients with PTH levels greater than 88 pmol/L confirmed on two laboratory tests for PTH taken at least 1 month apart.

**Exclusion Criteria**
Patients with primary hyperparathyroidism or parathyroid carcinoma.

**Initial Approval duration**

**Cinacalcet**
**Brand(s): Sensipar**
**DOSAGE FORM/ STRENGTH: 30 mg, 60 mg, 90 mg tablets**

Patients who are not a candidate for parathyroidectomy due to high surgical or anesthetic risk: 1 year

ii) Patients wait-listed for a parathyroidectomy requiring bridge therapy with Sensipar or awaiting an imminent renal transplant will be approved to the estimated date of the surgery.

**Duration of Approval:** 1 year or to the estimated date of the procedure for those using for bridge therapy and awaiting surgery.

**Renewals** will be considered for patients who are not candidates for parathyroidectomy and who continue to benefit from therapy. Requests for renewals should include the patient’s PTH level.

**Renewals** will NOT be considered for patients who have had a parathyroidectomy.

**Duration of Approval:** 1 year

**Sodium Thiosulfate**
**Brand(s): Seacalphyx DIN 02386666 and Hope pharmaceuticals DIN 02428393**
**DOSAGE FORM/ STRENGTH: 250mg/mL Injection; 12.5 g/50 mL Injection**
**Effective date: March 28, 2017**

**Approval of sodium thiosulfate for the treatment of calciphylaxis** will be provided where all of the following criteria have been met:

Patients with G4 or G5 chronic kidney disease; **AND**

i) Have been diagnosed with calciphylaxis either by:

a. 99m Technicium scintigraphy (bone scan) showing deposits that correspond to clinical lesions; OR

b. Biopsy; OR

c. Where scintigraphy negative and biopsy is not feasible, then diagnosis must be confirmed by nephrologist with submission of anonymized photographs of lesions AND a differential diagnosis checklist (e.g. warfarin-induced necrosis if on warfarin; lipohypertrophy if on insulin; cellulitis, nephrogenic sclerosing dermopathy, emboli, thrombi, fibrointimal hyperplasia and so on which depends on the site of lesion); **AND**
ii) Patient has either:
   a. Ulcerated lesions; OR

**Sodium thiosulfate**  
*Brand(s): Seacalphyx DIN 02386666 and Hope pharmaceuticals DIN 02428393*  
*DOSAGE FORM/ STRENGTH: 250mg/mL Injection, 12.5 g/50 mL Injection*

b. Non-ulcerated lesions which have not improved after 2 weeks of multimodal treatment with replacement of calcium-containing phosphate binders with non-calcium containing binders (i.e. sevelamer), discontinuation of vitamin D analogs and initiation of calcimimetic (i.e. cinacalcet), changes in dialysis prescription (reduction in dialysate calcium; consideration of increased dialysis intensity), replacement of warfarin with alternative anticoagulants where possible, wound management strategies, and analgesia for lesion pain.

**Requests for patients with calciphylaxis who do not meet the above criteria will be considered on a case-by-case basis.**

**Duration of Approval:** 2 months

**Renewals** will be considered for patients responding to treatment with improved pain control AND reduction in lesion number or size, reduction in ulcer size, or complete ulcer healing.

**Recommended dose:** 25g three times weekly.

**Duration of Approval:** Two months at a time until lesions are completely resolved, and for additional 2 months after complete healing.
Febuxostat
Brand(s): Uloric and generics see formulary OFIs
DOSAGE FORM/ STRENGTH: 80 mg

For the treatment of patients with documented severe allopurinol hypersensitivity syndrome* where lowering uric acid is recommended by clinical practice guidelines,

OR

For the treatment of patients with recurrent gout attacks despite treatment with allopurinol at a dose of 300 mg or more per day for at least six (6) months.

* For the purpose of the criteria severe allopurinol hypersensitivity syndrome is defined as follows;

The patient has had a clear exposure to allopurinol and:

(a) at least TWO of the following major clinical criteria:
   i) worsening renal function;
   ii) acute hepatocellular injury;
   iii) a rash including either toxic epidermal necrolysis (“TEN”), Stevens-Johnson syndrome (“SJS”), erythemamultiforme, generalised maculopapular exanthema or generalized exfoliative dermatitis (“GED”);

OR

(b) one of the major clinical criteria listed above in (a) and at least ONE of the following minor criteria:
   (i) fever
   (ii) eosinophilia
   (iii) leukocytosis

Note that an intolerance to allopurinol that does not meet the above criteria will not be eligible for reimbursement.

Duration of Approval: 1 year

Renewals will be considered in patients with objective evidence to demonstrate a benefit from treatment, documented as either a reduction in gout attacks or a reduction in uric acid levels.

Duration of Approval: 5 years
GRANULOMATOSIS WITH POLYANGIITIS OR MICROSCOPIC POLYANGIITIS

Rituximab -See Formulary for funded biosimilars
Brand(s): Riximyo, Ruxience, Truxima (biosimilars); Rituxan (Originator)
DOSAGE FORM/ STRENGTH: 10 mg/mL intravenous injection

For the induction of remission of severely active Granulomatosis with Polyangiitis (GPA) OR microscopic polyangiitis (MPA) as combination treatment with glucocorticoids, in patients who meet all of the following criteria:

1. The patient must have severe active disease that is life- or organ-threatening. At least one supporting laboratory and/or imaging report must be provided. The organ(s) and how the organ(s) is (are) threatened must be specified.

2. There is a positive serum assays for either proteinase 3-ANCA (anti-neutrophil cytoplasmic autoantibodies) or myeloperoxidase-ANCA. A copy of the laboratory report must be provided.

3. Cyclophosphamide cannot be used for the patient for at least ONE of the following reasons:
   i) The patient has failed a minimum of six IV pulses of cyclophosphamide; OR
   ii) The patient has failed three months of oral cyclophosphamide therapy; OR
   iii) The patient has a severe intolerance or an allergy to cyclophosphamide; OR
   iv) Cyclophosphamide is contraindicated; OR
   v) The patient has received a cumulative lifetime dose of at least 25 g of cyclophosphamide; OR
   vi) The patient wishes to preserve ovarian/testicular function for fertility.

Rituximab
Brand(s): Riximyo, Ruxience, Truxima (Biosimilars; Rituxan (Originator)
DOSAGE FORM/ STRENGTH: 10 mg/mL intravenous injection

The initial treatment would be a once weekly infusion dosed at 375 mg/m² x 4 weeks. The physician must confirm that the treatment would not be a maintenance infusion as maintenance infusions will not be funded.

Renewals will be considered provided that, the patient meets the same criteria for initial approval and the request for retreatment is made no less than 6 months after the last dose of the patient’s last treatment cycle with rituximab.

First Renewal: 1 year

Subsequent Renewals after first renewal: 2 years

(Rituximab is funded for course of therapy to be given at an interval of at least 6 months)
Rituximab will be funded as maintenance therapy for patients with severely active Granulomatosi
s with Polyangiitis [(GPA), also known as Wegner’s Granulomatosis (WG)] OR microscopic polyangii
ts (MPA). Patient must meet all of the following criteria:

a) The patient must have severe active disease that is life- or organ-threatening. At least one supporting laboratory and/or imaging report must be provided. The organ(s) and how the organ(s) is(are) threatened must be specified.
b) There is a positive serum assay for either proteinase 3-ANCA (anti-neutrophil cytoplasmic autoantibodies) or myeloperoxidase-ANCA. A copy of the laboratory report must be provided.
c) Stabilization of the condition with induction doses of cyclophosphamide (IV or PO doses) and a glucocorticoid as combination over 4 to 6 months until disease remission followed by rituximab at 500mg doses every 6 months. Cyclophosphamide dosing to align with MAINRITSAN studies1.

After remission (typically within a month of remission), rituximab will be administered as one of the following:

1. A fixed dose regimen of Rituximab consisting of 500 mg dosed at days 0 and 14 followed by fixed doses of 500 mg at 6, 12, and 18 months, used in combination with low-dose prednisone of another glucocorticoid; x 18 months duration of funding OR
2. A tailored dose regimen of Rituximab based on CD19 and ANCA monitoring. Dose of Rituximab funded is 500 mg on day 0 followed by a dose as early as every 3 months if CD19 exceeds 0/mm3 or if ANCA reappears or if there is a titre increase x 18 months duration of funding.

1Remission-induction therapy included prednisone (starting at 1 mg per kilogram of body weight per day, followed by gradual tapering), preceded in some patients by methylprednisolone “pulses” (500 to 1000 mg daily for 1 to 3 consecutive days), and

**Rituximab**

**Brand(s):** Riximyo, Ruxience, Truxima (Biosimilars; Rituxan (Originator))

**DOSAGE FORM/ STRENGTH:** 10 mg/mL intravenous injection

“pulse” cyclophosphamide (0.6 g per square meter of body-surface area on days 0, 14, and 28, then 0.7 g per square meter every 3 weeks for three to six additional pulses) until remission was attained, after 4 to 6 months. At that time, and within a maximum of 1 month after the last cyclophosphamide pulse, we have also accepted oral dosing (an example of oral cyclophosphamide dosing that has been used by clinicians is 150 mg daily).

Approval duration: 18 months

Renewals:
Renewals will be considered case-by-case. Requests should include information pertaining to the number of disease flares during the period of funding and a description of symptoms during flares. The dosing interval of use must be maintained as every 6 months and should be specified.

**HEPATOLOGY DRUGS**

**HEPATITIS C DRUGS**

The following drugs are reimbursed on the Ontario drug benefit formulary as limited use benefits for patients with Chronic Hepatitis C Infection upon meeting the LU criteria:

i) Epclusa (sofosbuvir / velpatasvir) 400mg/100mg Tab

ii) Harvoni (ledipasvir / sofosbuvir) 90mg/400mg Tab (GIL)

iii) Ibavir (ribavirin) 200mg, 400mg, 600mg Tab

iv) Maviret (glecaprevir/pibrentasvir) 100 mg/40 mg Tab

v) Sovaldi (sofosbuvir) 400mg Tab

vi) Vosevi (sofosbuvir/velpatasvir/voxilaprevir) 400 mg/100 mg/100 mg Tab

vii) Zepatier (elbasvir / grazoprevir) 50mg/100mg Tab

The Ministry only considers funding of patient with Chronic Hepatitis C infection.

Please refer to the Limited Use Criteria in the Ontario Drug Benefit Formulary for provincial reimbursement criteria for these products which are part of Ontario’s hepatitis C framework.

Patients not meeting limited use criteria may be considered on a case-by-case basis through the Exceptional Access Program.

**Adefovir**

Brand(s): Hepsera

**DOSAGE FORM/ STRENGTH:** 10 mg tablet

For the treatment of chronic hepatitis B in patients with objective evidence of lamivudine virologic* breakthrough where failure is not due to poor adherence to therapy; AND

- Liver biopsy showing Metavir stage 3 fibrosis or greater; OR
- Documented evidence of cirrhosis
OR

- Patients with the presence of a lamivudine resistance mutation****; AND
- Liver Biopsy showing Metavir stage 3 fibrosis or greater; OR
- Documented evidence of cirrhosis

**Duration of Approval:** 1 year (If Cirrhotic: Lifetime)

Duration of Approval for Renewal: 5 years

**Note:** Effective February 28, 2018, Entecavir, Lamivudine, and Tenofovir became a Limited Use Benefit on the Ontario Drug Benefit Formulary – Please refer to the formulary for the Limited Use Criteria.

**Interferon – alpha-2b**
Brand(s): Intron A
**DOSAGE FORM/ STRENGTH:** 18 MU, 30 MU, 60 MU; 18 MU/3 mL, 10 MU/mL, 25 MU/2.5 mL vials

For the treatment of chronic hepatitis B where the patient meets the following criteria:

- Patients less than 50 years of age; AND
- 2 ALTs > 2 x ULN within the past 6 month period; AND
- HBV DNA between $1 \times 10^4 - 1 \times 10^7$ IU/mL; AND
- Metavir stage 3 fibrosis or less (i.e. no cirrhosis)

Requests for pediatric patients will be considered case-by-case.

**Duration of Approval:** HBeAg pos: 24 weeks, HBeAg neg: 48 weeks
Obeticholic Acid (Ocaliva) will be funded for the treatment of primary biliary cholangitis (PBC) in adult patients who meet the following criteria:

i) Diagnosis of PBC is demonstrated by antimitochondrial antibodies or a liver biopsy; AND

ii) Used in combination therapy with ursodeoxycholic acid (UDCA) in patients who have experienced an inadequate response\(^1\) to a minimum of twelve months of treatment with UDCA OR as monotherapy in patients who have experienced unmanageable intolerance to UDCA; AND

iii) The request is prescribed by or in consultation with a prescriber who is a gastroenterologist, hepatologist or internist with experience in the treatment of PBC. (If you are a prescriber who is not one of the specialists identified above, please submit the consultation note with the request.)

\(^1\)Note that an inadequate response is defined as a patient who has used UDCA to treat PBC for a minimum of twelve (12) months and demonstrates ANY ONE or more of the following:

a) alkaline phosphatase $\geq 1.67 \times$ upper limit of normal

b) total bilirubin $> 1 \times$ upper limit of normal and $< 2 \times$ upper limit of normal

c) abnormal bilirubin with progressing and/or compensated cirrhosis

(Documentation of lab work to be submitted with the request application.)

Renewals will be considered in patients who continue to benefit from treatment as evidenced by any one of the following:

(a) a reduction in the alkaline phosphatase level to less than $1.67 \times$ upper limit of normal; and/or

(b) a 15% reduction in the alkaline phosphatase level compared with baseline values prior to initiation of treatment with obeticholic acid; and/or

(c) a normal bilirubin level.

and the patient has not developed unacceptable toxicity from treatment with obeticholic acid.
Obeticholic Acid
Brand(s): Ocaliva
DOSAGE FORM/ STRENGTH: 5mg, 10mg tablet

Patients not meeting the above renewal criteria may be considered on a case-by-case basis.

Exclusion Criteria (applies to both initial and renewal requests):

Pre-liver transplant patients and/or patients who have complete biliary obstruction will not be funded.

Duration of funding approval for initials: 1 year

Duration of funding approval for renewals: 1 year
INFLAMMATORY BOWEL DISEASES

Adalimumab – See Formulary for funded biosimilars
Brand(s): Humira and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 40 mg/0.8 mL prefilled syringe, 40mg/0.8mL and 20 mg/0.2 mL prefilled pens for subcutaneous injection

For the treatment of fistulising Crohn’s disease with concomitant luminal disease in patients who meet the following criteria;

- Patient with actively draining perianal or enterocutaneous fistula(e) that have recurred or persist despite a course of appropriate antibiotic therapy (e.g. ciprofloxacin and/or metronidazole) AND immunosuppressive therapy (e.g. azathioprine or 6-mercaptopurine) AND

- Harvey Bradshaw Index (HBI) score ≥ 7

The dose that will be considered is Adalimumab 160 mg at week zero, 80 mg at week two, followed by 40 mg every two weeks.

Duration of Approval: 3 months

Renewal will be considered based on the response to therapy.

The dose that will be considered on renewals is Adalimumab 40 mg every two weeks. All requests for higher doses will not be approved.

Duration of Approval: 3 months to 1 year pending fistula(e) resolution

Second Renewal:

2 years for 2nd renewal of requests with complete resolution

Case-by-case duration for renewal of requests with partial resolution

Pediatric patients will be considered case-by-case.
Treatment of **moderate to severe** (luminal) Crohn’s Disease in patients who have:

- HBI (Harvey Bradshaw Index) score ≥7*; and
- Failed to respond to conventional treatment with glucocorticoids (prednisone 40mg/day or equivalent for at least 2 weeks or dose cannot be tapered to below prednisone 20 mg/day or equivalent); and
- Failed to respond to an immunosuppressive agent (azathioprine, 6-mercaptopurine, methotrexate, or cyclosporine) tried for at least 3 months.

*If the patient has HBI <7, the request will be reviewed by external medical experts when the following information is provided: bloodwork (with hematocrit, hemoglobin, C reactive protein, ESR, platelets, and ferritin levels); supporting endoscopy; details of weight loss; and a list of narcotic analgesics being used.

Pediatric patients will be considered case-by-case.

**Duration of Approval:** 6 months

**Renewal** will be considered for patients with 50% reduction in HBI from pre-treatment as well as improvement of symptoms (e.g., absence of bloody diarrhea and weight stabilization or increase) and no longer using steroids. Biochemical improvements may also be required.

The planned dosing regimen for the requested biologic should be provided.

The recommended: Adalimumab: 160mg at week 0; 80mg at week 2; followed by 40mg every two weeks

**Duration of Approval:** First renewal: 1 year

Second and subsequent renewals: 2 years
Adalimumab – See Formulary for funded biosimilars
Brand(s): Humira and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 40 mg/0.8 mL prefilled syringe, 40mg/0.8mL and 20 mg/0.2 mL prefilled pens for subcutaneous injection

For the treatment of ulcerative colitis disease in adult patients\(^1\) who meet the following criteria: Induction Criteria
Mild disease
  a. Mayo score <6 AND
  b. Patients with mild disease will be considered on a case-by-case basis BUT
     submission must include the rationale for coverage
Moderate disease
  a. Mayo score between 6 and 10 (inclusive); AND
  b. Endoscopic* subscore of 2; AND
  c. Failed 2 weeks of oral prednisone at daily doses ≥40mg (or a 1 week course of IV equivalent) AND 3 months of azathioprine (AZA)/ 6-mercaptopurine (6MP) (or where the use of immunosuppressants is contraindicated):
     OR
     Stabilized with 2 weeks of oral prednisone at daily dose ≥ 40mg (or a 1 week course of IV equivalent) but the prednisone dose cannot be tapered despite 3 months of AZA/ 6MP (or where the use of immunosuppressants is contraindicated)
Severe disease
  a. Mayo score >10 AND
  b. Endoscopic* subscore of ≥2 AND
  c. Failed 2 weeks of oral prednisone at daily dose ≥ 40mg (or 1 week IV equivalent); OR
     Stabilized with 2 weeks oral prednisone ≥ 40mg (or 1 week of IV equivalent) but the prednisone dose cannot be tapered despite 3 months of AZA/6MP (or where the use of immunosuppressants is contraindicated)

AZA/6MP (or where the use of immunosuppressants is contraindicated)
  Initial Approval: 6 months at 160 mg initially administered at week 0, followed by 80mg at week 2, then 40 mg every other week thereafter.

*The endoscopy procedure must be done within the
  12 months prior to initiation of treatment
Maintenance Criteria
After 8 weeks of adalimumab therapy:
  a. Mayo score <6 AND
  b. 50% reduction in prednisone from the starting dose
Approval: 6 months at 40mg every other week.
If patient is completely off steroids,
Approval: 12 months at 40 mg every other week.
Subsequent renewals:
  a. Mayo score <6; AND
  b. Must be completely off steroids
Approval: 2 years at 40mg every other week.
(Patients who remain on steroids will be considered on a case-by-case basis)
1Pediatric patients will be considered case-by-case.
Infliximab—See formulary for funded biosimilars
Brand(s): Avsola, Inflectra, Renflexis (Biosimilars); Remicade (Originator)
DOSAGE FORM/STRENGTH: 100mg/vial Injection for Infusion

Inflectra (Since November 30, 2016), Renflexis (Since September 27, 2018) and Avsola (since December 18, 2020) are biosimilars that are publicly funded for the treatment of patients with Inflammatory Bowel Diseases (i.e. Crohn’s Disease or Ulcerative Colitis) upon meeting the Limited Use Criteria on the Ontario Drug Benefit (ODB) formulary. Requests for Inflectra, Renflexis, or Avsola not meeting the LU criteria may be forwarded to EAP for case-by-case assessment.

Requests for infliximab in treatment-naïve patients with Crohn’s Disease or Ulcerative Colitis will be required to consider treatment using one of the biosimilars (Inflectra or Renflexis or Avsola). Patients initiated on Remicade by the manufacturer’s patient support program after the effective date of public funding of the biosimilar infliximab products may be expected to provide ongoing access to their patient.

Treatment of moderate to severe (luminal) Crohn’s Disease in patients who have:

- HBI (Harvey Bradshaw Index) score ≥7*; and
- Failed to respond to conventional treatment with glucocorticoids (prednisone 40mg/day or equivalent for at least 2 weeks or dose cannot be tapered to below prednisone 20 mg/day or equivalent); and
- Failed to respond to an immunosuppressive agent (azathioprine, 6-mercaptopurine, methotrexate, or cyclosporine) tried for at least 3 months.

Note: Any intolerance(s) or contraindication(s) to treatment with required alternative(s) must be described in detail.

*If the patient has HBI <7, the request will be reviewed by external medical experts when the following information is provided: bloodwork (with hematocrit, hemoglobin, C reactive protein, ESR, platelets, and ferritin levels); supporting endoscopy; details of weight loss; and a list of narcotic analgesics being used.

Pediatric patients will be considered case-by-case.

Duration of Approval: 6 months

Renewal will be considered for patients with 50% reduction in HBI from pre-treatment as well as improvement of symptoms (e.g., absence of bloody diarrhea and weight stabilization or increase) and no longer using steroids. Biochemical improvements may also be required.

The planned dosing regimen for the requested biologic should be provided.
Infliximab – See Formulary funded biosimilars
Brand(s): Avsola, Inflectra, Renflexis (Biosimilars); Remicade (Originator)
DOSAGE FORM/ STRENGTH: 100mg/vial Injection for infusion

**Recommended dose:** Infliximab 5 mg/kg/dose at 0, 2 and 6 weeks followed by 5mg/kg/dose every 8 weeks

Requests for higher doses of infliximab must provide a description of symptoms and HBI score on standard dosing and may include laboratory support of infliximab levels for consideration of case-by-case consideration.

**Duration of Approval:** First renewal: 1 year
Second and subsequent renewals: 2 years

**Remicade for fistulizing Crohn’s disease:**
Actively draining perianal or enterocutaneous fistula(e) that have recurred or persist despite a course of:
- appropriate antibiotic therapy (e.g., ciprofloxacin and/or metronidazole); AND
- immunosuppressive therapy (azathioprine or 6-mercaptopurine therapy).

Duration of approval: Infliximab 3 doses of 5mg/kg/dose, administered at 0, 2 and 6 weeks.

**Duration of Approval:** 6 months

If the patient has been using a higher dosing regimen over the past year, the requesting MD must provide the rationale for this dose by comparing the patient’s symptoms on standard dosing and the current dosing. Then the request should be sent for external review.

**Renewal of funding of patients using Remicade for the treatment of fistulizing Crohn’s Disease** will be considered for patients with resolution of fistulae.

The planned dosing regimen for the requested biologic should be provided. The recommended dose for the treatment of Crohn’s Disease is 5 mg/kg/dose at 0, 2 and 6 weeks followed by 5mg/kg/dose every 8 weeks with up to 10 mg/kg/dose every 8 weeks being considered on a case-by-case basis.

**Approval duration of first renewal:** 6 months to 1 year pending fistula(e) resolution

**Approval duration of second and subsequent renewals:** 2 years with complete resolution; case-by-case duration with partial resolution
Infliximab – See Formulary funded biosimilars
Brand(s): Avsola, Inflectra, Renflexis (Biosimilars); Remicade (Originator)
DOSAGE FORM/ STRENGTH: 100mg/vial Injection for infusion

Initial induction requests for infliximab for patients with mild Ulcerative Colitis (Mayo score < 6) may be considered for Infliximab as Inflectra on a case-by-case basis through EAP but the submission must include the rationale for coverage.

Patients treatment experienced to Remicade and transitioning to public funding must meet the initiation (induction) criteria before consideration of funding of maintenance under renewal criteria will be applied.

Induction (Initiation) Criteria

1. Moderate disease
   - Mayo score between 6 and 10 (inclusive); AND
   - *Endoscopic subscore of 2; AND
   - Failed 2 weeks of oral prednisone ≥ 40mg (or IV equivalent for at least 1 week) AND
     3 months of azathioprine(AZA)/ 6-mercaptopurine (6-MP) (or where the use of immunosuppressants is contraindicated**)
     OR
     Stabilized with 2 weeks of oral prednisone ≥40mg (or a 1 week course of IV equivalent) but the prednisone dose cannot be tapered despite 3 months of AZA/6MP (or where the use of immunosuppressants is contraindicated**)

*The endoscopy procedure must be done within the last year but does not have to be full endoscopy.

**Contraindication to Aza/6MP includes pancreatitis, allergic reaction [fever and/or rash and arthritis], malaise, diarrhea and hepatitis

Approval duration: 6 months

Approved Dose: Infliximab 5mg/kg/dose at 0, 2 and 6 weeks followed by 5mg/kg/dose every 8 weeks.

2. Severe disease
   - Mayo score >10; AND
   - *Endoscopy subscore of 2 or more; AND
   - Failed 2 weeks of oral prednisone ≥40mg (or 1 week IV equivalent)
     OR
     Stabilized with 2 weeks of oral prednisone ≥40mg (or 1 week of IV equivalent) but the prednisone dose cannot be tapered despite 3 months of Aza/6MP (or where the use of immunosuppressants is contraindicated**)
**Infliximab – See Formulary funded biosimilars**
Brand(s): Avsola, Inflectra, Renflexis (Biosimilars); Remicade (Originator)

**DOSAGE FORM/ STRENGTH:** 100mg/vial Injection for infusion

*The endoscopy procedure must be done within the last year but does not have to be full endoscopy.

**Contraindication to Aza/6MP includes pancreatitis, allergic reaction [fever and/or rash and arthritis], malaise, diarrhea and hepatitis

Approval duration: 6 months
Dose: Remicade 5mg/kg/dose at 0, 2 and 6 weeks followed by 5mg/kg/dose every 8 weeks.

**Maintenance (Renewal) Criteria for first renewal**

After 3 loading doses of Remicade if Mayo score <6 AND 50% reduction in prednisone from the starting dose

Approval duration: 6 months
If After 3 loading doses of Remicade if Mayo score <6 AND patient is no longer on prednisone

Approval duration: 12 months
Approved Dose: Infliximab 5mg/kg/dose up to every 6 weeks

**Maintenance (Renewal) Criteria for second and subsequent renewals**

a. Mayo score <6* AND  
b. Must be off steroids

Patients who remain on steroids will be considered on a case-by-case basis.

Approval duration: 12 months to up to 2 years for those off steroids

Approved Dose: 5 mg/kg/dose up to every 6 weeks

1Note that the endoscopy procedure must be done within the last year but does not have to be full endoscopy.

**Pediatric patients will be considered case-by-case.**
Golimumab
Brand(s): Simponi
DOSAGE FORM/ STRENGTH: 50 mg/0.5mL Pre-Filled Syringe Or Auto-Injector, 100 mg/ mL Pre-filled Syringe or Auto-Injector

For the treatment of ulcerative colitis disease in patients who meet the following criteria:

**Induction Criteria**

Mild disease
- a. Mayo score <6 AND
- b. Patients with mild disease will be considered on a case-by-case basis BUT submission must include the rationale for coverage

Moderate disease
- a. Mayo score between 6 and 10 (inclusive) AND
- b. Endoscopic subscore of 2 AND
- c. Failed 2 weeks of oral prednisone ≥40mg (or a 1 week course of IV equivalent) but the prednisone dose cannot be tapered despite 3 months azathioprine(AZA)/6-mercaptopurine(6MP) (or where the use of immunosuppressants is contraindicated)

        OR

        Stabilized with 2 weeks of oral prednisone ≥ 40mg (or a 1 week course of IV equivalent) but the prednisone dose cannot be tapered despite 3 months of AZA/6MP (or where the use of immunosuppressants is contraindicated)

Severe disease
- a. Mayo score >10 AND
- b. Endoscopy subscore of ≥2 AND
- c. Failed 2 weeks of oral prednisone ≥ 40mg (or 1 week IV equivalent)

        OR

        d. Stabilized with 2 weeks oral prednisone ≥ 40mg (or 1 week of IV equivalent) but the prednisone dose cannot be tapered despite 3 months of AZA/6MP (or where the use of immunosuppressants is contraindicated)
Golimumab
Brand(s): Simponi
DOSAGE FORM/ STRENGTH: 50 mg/0.5mL Pre-Filled Syringe Or Auto-Injector, 100 mg/ mL Pre-filled Syringe or Auto-Injector

Initial Approval: 6 months at 200 mg initially administered at week 0, followed by 100mg at week 2, and then 50 mg every 4 weeks thereafter. The maintenance dose of 100mg every 4 weeks can be considered at the discretion of the treating physician

Maintenance Criteria

After 4 loading doses of Simponi:

a. Mayo score <6 AND
b. 50% reduction in prednisone from the starting dose

Approval: 6 months at 50 mg or 100 mg every 4 weeks

If patient is completely off steroids.

Approval: 12 months at 50 mg or 100 mg every 4 weeks.

Subsequent renewals:

a. Mayo score <6; AND
b. Must be off steroids

(Patients who remain on steroids will be considered on a case-by-case basis)

Approval: 2 years at 50 mg or 100 mg every 4 weeks.

Duration of Approval: Renewal duration: 6 months to 1 year (Pending if patient continues on steroids.)

Second and subsequent renewal 2 years for those off steroids:
Applications for reimbursement should be accompanied by a copy of the parenteral support weekly usage volumes at baseline prior to start of teduglutide and must include the parenteral support usage volumes for the 4 weeks of use prior to the date that the Ministry funding for initial or renewal is requested.

Adults (age ≥ 18 years) - Initial Criteria:

For the ongoing funding of patients who have turned 18 years of age who were previously funded upon meeting EAP funding criteria and who continue to meet the adult renewal criteria; OR

For the ongoing treatment of new patients requesting public funding who are at least 18 years of age with Short Bowel Syndrome (SBS) who meet the following criteria:

Prior to starting teduglutide (Revestive), the patient meets the following;

- Has short bowel syndrome (SBS) as a result of major intestinal resection due to injury, volvulus, vascular disease, cancer, or Crohn’s Disease\(^1\); AND
- Patient’s intestinal resection has resulted in dependency on parenteral nutrition for at least 12 months; AND
- Patient requires parenteral nutrition required at least three times weekly to meet caloric, fluid or electrolyte needs due to ongoing malabsorption; AND
- Patient’s frequency and volume of parenteral nutrition has been stable for at least one month;

Those who are initiated on teduglutide upon meeting the above criteria will be reimbursed for ongoing access of teduglutide through the Exceptional Access Program (EAP) upon meeting the below criteria;

- The patient has achieved at least a 20% reduction in Parenteral Support volume while on teduglutide in the 4 weeks prior to the application for funding to EAP compared to their baseline measures in the 4 weeks prior to start of teduglutide.\(^2\)\(^3\)

\(^1\) Case-by-case consideration will be provided for requests for patients with short bowel syndrome not due to the reasons provided.

\(^2\) Parenteral Support volumes measurements are to compare the 4 weeks of use prior to start of teduglutide with the 4 weeks of use prior to the date that EAP request for funding is provided.

\(^3\) For adult patients (i.e. at least 18 years of age) requesting EAP funding for ongoing teduglutide therapy who initiated on teduglutide between 1 and 17 years of age that was
Teduglutide
Brand(s): Revestive
DOSAGE FORM/ STRENGTH: 5 mg vials for Injection

not publicly funded, ongoing response to teduglutide will be evaluated through comparing the patient’s current (i.e. while on teduglutide in the 4 weeks prior to the application for funding to EAP) Parenteral Support volume to the patient’s adjusted initial Parenteral Support volume; for clarity, adjusted initial Parenteral Support volume is defined as the patient’s last available weight as a pediatric multiplied by the patient’s baseline Parenteral Support volume and divided by the patient’s baseline weight at initiation (i.e. in the 4 weeks prior to start of teduglutide as a pediatric) of teduglutide.

Duration of approval: 12 months

Exclusion Criteria:
• Patients with active gastrointestinal malignancy or history of gastrointestinal malignancy in the past 5 years before start of treatment

Renewal Criteria:

Renewals will be considered in those who are able to demonstrate a reduction in their Parenteral Support volume requirements by at least 20% compared to baseline measures prior to commencing therapy with teduglutide. 3
Applications for renewal of reimbursement should be accompanied by a copy of the Parenteral Support weekly usage volumes for the 4 weeks of use prior to the date that the Ministry funding is requested.
Duration of approval: 12 months

Discontinuation/Stopping Criteria:
Patients will not be approved for funding by the EAP if a 20% reduction in the usual average weekly volume of Parenteral Support solutions prior to starting therapy with teduglutide has not been achieved/maintained by 52 weeks of treatment with teduglutide.
**Teduglutide**
*Brand(s): Revestive*

**DOSAGE FORM/ STRENGTH:** 5 mg vials for Injection

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**Pediatrics (age between 1 and 17 years) - Initial Criteria:**

For the ongoing treatment of patients between 1 and 17 years of age with Short Bowel Syndrome (SBS) who are dependent on Parenteral Support who meet the following criteria:

Prior to starting teduglutide (Revestive), the patient meets the following criteria:

- Has short bowel syndrome (SBS) as a result of major intestinal resection; AND
- Parenteral Support requirements must be stable or there must have been no improvement in enteral feeding for at least the preceding three months; AND
- The cumulative lifetime duration of Parenteral Support therapy must be at least 12 months; AND
- Patient requires parenteral support to provide more than 30% of caloric, and/or fluid/electrolyte needs due to ongoing malabsorption; AND
- Request is from a physician or nurse practitioner working within a specialized multi-disciplinary intestinal rehabilitation program.

Those who are initiated on teduglutide upon meeting the above criteria will be reimbursed for ongoing access of teduglutide through the Exceptional Access Program (EAP) upon meeting the below criteria:

- Initial treatment response is assessed 6 months or more after initiating treatment with teduglutide.
- The patient has achieved at least a 20% reduction in weight-adjusted Parenteral Support volume while on teduglutide in the 4 weeks prior to the application of funding to EAP compared to their baseline weight-adjusted Parenteral Support volume measures in the 4 weeks prior to start of teduglutide.\(^1\)\(^2\)

\(^1\) Weight-adjusted Parenteral Support volume is defined as the patient’s Parenteral Support volume divided by the Patient’s weight.

\(^2\) Parenteral Support volumes measurements are to compare the 4 weeks of use prior to start of teduglutide with the 4 weeks of use prior to the date that EAP request for funding is provided.
**Teduglutide**
Brand(s): Revestive
DOSAGE FORM/ STRENGTH: 5 mg vials for Injection

Notes:

a. Parenteral Support volume and percentage of total consumption should be documented at each clinic visit.

b. Assessments for subsequent renewals should be carried out at 6 months intervals.

Duration of first EAP approval: 7 months

**Exclusion Criteria:**

- Patients with active gastrointestinal malignancy or history of gastrointestinal malignancy in the past 5 years before start of treatment

**Renewal Criteria:**

Renewals will be considered in patients who are able to demonstrate a reduction in their weight-adjusted Parenteral Support volume requirements by at least 20% compared to baseline weight-adjusted Parenteral Support volume measures prior to commencing therapy with teduglutide.¹

Applications for renewal of reimbursement should be accompanied by a copy of the weight-adjusted Parenteral Support weekly usage volumes for the 4 weeks of use prior to the date that the Ministry funding is requested.

Duration of approval: 12 months

**Discontinuation/Stopping Criteria**

Patients meeting initial funding criteria will not be approved for ongoing funding by the EAP if a 20% reduction in the usual average weekly volume of weight-adjusted Parenteral Support solutions prior to starting therapy with teduglutide has not been achieved/maintained at the time of renewal.
Vedolizumab
Brand(s): Entyvio
DOSAGE FORM/ STRENGTH: 300 mg Injection

For the treatment of fistulising Crohn’s disease with concomitant luminal disease inpatients who meet the following criteria;

- Patient with actively draining perianal or enterocutaneous fistula(e) that have recurred or persist despite a course of appropriate antibiotic therapy (e.g. ciprofloxacin and/or metronidazole) AND immununosuppressive therapy (e.g. azathioprine or 6-mercaptopurine) AND
- Harvey Bradshaw Index (HBI) score ≥ 7

Initial Approval: 6 months at 300 mg initially administered at week 0, followed by 300mg at week 2, 300mg at week 6, then 300 mg every 8 weeks thereafter.

Renewal will be considered based on the response to therapy. The dose that will be considered on renewals is Vedolizumab (Entyvio) is 300 mg every eight weeks.

Approval duration:
First renewal: 6 months to 1 year pending fistula(e) resolution
Second and subsequent renewals: 2 years with complete resolution; case-by-case duration with partial resolution

For the treatment of moderate to severe (luminal) Crohn’s Disease in patients who have:
- HBI (Harvey Bradshaw Index) score ≥7*; AND
- Failed to respond to conventional treatment with glucocorticoids (prednisone 40mg/day or equivalent for at least 2 weeks or dose cannot be tapered to below prednisone 20 mg/day or equivalent); AND
- Failed to respond to an immunosuppressive agent (azathioprine, 6-mercaptopurine, methotrexate, or cyclosporine) tried for at least 3 months.

Note: Any intolerance(s) or contraindication(s) to treatment with required alternative(s) must be described in detail.
**Vedolizumab**

**Brand(s):** Entyvio  
**DOSAGE FORM/ STRENGTH:** 300 mg Injection

*If the patient has HBI <7, the request will be reviewed by external medical experts when the following information is provided: bloodwork (with hematocrit, hemoglobin, C reactive protein, ESR, platelets, and ferritin levels); supporting endoscopy; details of weight loss; and a list of narcotic analgesics being used.

*Initial Approval: 6 months at 300 mg initially administered at week 0, followed by 300mg at week 2, 300mg at week 6, then 300 mg every 8 weeks thereafter.*

**Renewal** will be considered for patients with 50% reduction in HBI from pre-treatment as well as improvement of symptoms (e.g., absence of bloody diarrhea and weight stabilization or increase) and no longer using steroids. Biochemical improvements may also be required.

**For the treatment of ulcerative colitis disease** in patients who meet the following criteria:

**Induction Criteria**

Mild disease

a. Mayo score <6 AND

b. Patients with mild disease will be considered on a case-by-case basis BUT submission must include the rationale for coverage

Moderate disease

a. Mayo score between 6 and 10 (inclusive) AND

b. Endoscopic* subscore of 2 AND

c. Failed 2 weeks of oral prednisone at daily doses ≥40mg (or a 1 week course of IV equivalent) and 3 months of azathioprine (AZA)/ 6-mercaptopurine (6MP) (or where the use of immunosuppressants is contraindicated)

    **OR**

Stabilized with 2 weeks of oral prednisone at daily dose ≥ 40mg (or a 1 week course of IV equivalent) but the prednisone dose cannot be tapered despite 3 months of AZA/ 6MP (or where the use of immunosuppressants is contraindicated)
Vedolizumab
Brand(s): Entyvio
DOSAGE FORM/ STRENGTH: 300 mg Injection

Severe disease

a. Mayo score >10 AND

b. Endoscopic* subscore of ≥2 AND

c. Failed 2 weeks of oral prednisone at daily dose ≥ 40mg (or 1 week IV equivalent)

OR

Stabilized with 2 weeks oral prednisone ≥ 40mg (or 1 week of IV equivalent) but the prednisone dose cannot be tapered despite 3 months of AZA/6MP (or where the use of immunosuppressants is contraindicated)

Initial Approval: 6 months at 300 mg initially administered at week 0, followed by 300mg at week 2, 300mg

at week 6, then 300 mg every 8 weeks thereafter.

*The endoscopy procedure must be done within the 12 months prior to initiation of treatment

Maintenance Criteria

After 3 loading doses of Entyvio:

a. Mayo score <6 AND

b. 50% reduction in prednisone from the starting dose

Approval: 6 months at 300 mg every 8 weeks

If patient is completely off steroids, Approval: 12 months at 300 mg every 8 weeks.

Subsequent renewals:

a. Mayo score <6; AND

b. Must be completely off steroids

Approval: 2 years at 300 mg every 8 weeks.

(Patients who remain on steroids will be considered on a case-by-case basis)
Zopiclone
Brand(s): Imovane + generic brands
DOSAGE FORM/ STRENGTH: 5 mg, 7.5 mg tablet

For the treatment of insomnia as a single hypnotic agent in patients who meet the following criteria;
  • Have failed at least two benzodiazepines; OR
  • Have failed or experienced intolerance to at least one benzodiazepine and one other hypnotic (i.e., amitriptyline, trazodone, etc)

Duration of Approval: 2 Years
For the treatment of insomnia if patient has an identified psychiatric diagnosis. Renewals will be considered in patients who are responding to therapy AND who continues to require therapy AND who are using zopiclone as a single agent.
Duration of Approval: 2 Years
METABOLIC/GENETIC MODIFIERS

Asfotase alfa
Brand(s): Strensiq
DOSAGE FORM/ STRENGTH: 18mg/0.45mL, 28mg/0.7mL, 40mg/1mL, 80mg/0.8mL

Initiation Criteria:

For the treatment of perinatal/infantile, childhood, or juvenile-onset hypophosphatasia (HPP) in patients who meet the following criteria:

- Diagnosis is confirmed by genetic testing (i.e. documented tissue-nonspecific alkaline phosphatase gene mutations); AND
- Serum alkaline phosphatase (ALP) level is below the age-adjusted normal range\(^1\) using age and gender adjusted norms; AND
- Plasma pyridoxal-5-phosphate (PLP) above the upper limit of normal; AND
- Radiologically confirmed HPP-related skeletal abnormalities; AND
- Diagnosis occurred before the patient’s 12th birthday with documented onset of signs/symptoms\(^2\) of HPP prior to their 12th birthday; AND
- Patient is younger than 18 years of age at the time the treatment is initiated; AND
- Patient does not have odonto- or pseudo-HPP (i.e. craniosynostosis alone, premature loss of deciduous teeth alone and vitamin D deficiency to be ruled out); AND
- The patient’s treatment plan and goals of therapy is provided prior to the initiation of therapy; AND
- Patient is under the care of a metabolic specialist with expertise in the diagnosis and management of HPP.

Approval duration for initial requests: 6 months

Renewal Criteria:

Renewals of reimbursement will be considered in patients meeting the following criteria:

- Patient continues to be under the care of a metabolic specialist; AND
- Patient has demonstrated compliance to the treatment and monitoring schedule: AND
- Pre-specified goals\(^3\) based on the patient’s clinical status at initiation of treatment are met and the patient is deemed to continue to benefit from treatment. (Note: The request must include information about the treatment responses and milestones)

First renewal: 6 months

Subsequent renewals: 1 year
Asfotase alfa
Brand(s): Strensiq
DOSAGE FORM/ STRENGTH: 18mg/0.45mL, 28mg/0.7mL, 40mg/1mL, 80mg/0.8mL

Stopping Criteria:
- Discontinuation to be considered after growth is completed based on objective measure of height, weight and closure of bone growth plates as confirmed radiologically.
- Babies with perinatal/infantile HPP who fail treatment trials of 6 months
- If pre-specified goals are not met at reassessment, the treatment should not be continued.

Notes:
1 Normal range as informed by the Canadian Laboratory Initiative on Paediatric Reference Intervals (CALIPER) can be used as a reference for this information. Below upper limit of normal refers to 2 or lower standard deviations above the mean.
2 Incoming Requests should address the following:
   - baseline skeletal symptoms including age and dates of for those assessments
   - abnormalities of skeletal mineralization
   - fracture history
   - growth plate irregularities and bone and skeletal growth
   - description of growth and developmental milestones
   - Signs, symptoms, and history of seizures
   - respiratory function including need for ventilator support
   - Activity and mobility
   - Laboratory markers that include vitamin D levels, calcium levels

Assessments such as the Radiographic Global Impressive of Change (RGIC) score and/or the Thacher score for evaluating rickets may be provided at baseline and at the time of renewal of coverage (as applicable) as a measure of response and benefit from therapy.

3 Specific patient treatment goals should be developed on a case-by-case basis and may include some of the following: Healing of rickets, improved bone mineralization, fewer fractures, reduced pain, improved growth, mobility, improvement in respiratory status, attainment of age-appropriate growth milestones, improvement in gait or deformities, improved quality of life measures.

Documentation of improvement from baseline is to be provided at the time of renewal.
Please note that effective April 8, 2021, foreign labelled Strensiq (generic name: asfotase alfa) has been added as a temporary Exceptional Access Program benefit due to drug shortage.

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<th>Brand Name</th>
<th>Generic Name</th>
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Initiation Criteria:

For the treatment of X-linked hypophosphatemia (XLH) in patients who meet all of the following criteria:

1. Treatment is initiated in a pediatric patient who is at least 1 year of age and in whom the epiphyseal closure has not occurred yet; AND
2. XLH diagnosis is confirmed by genetic documentation of 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; AND
3. Clinical presentation consistent with XLH diagnosis, including fasting hypophosphatemia\(^1\) and normal renal function (defined as fasting serum creatinine level below the age-adjusted upper limit of normal); AND
4. Radiographic evidence of rickets with a baseline\(^2\) rickets severity score (RSS) total score of 2 or greater at the time Crysvita treatment is initiated; AND
5. Patient is under the care of a specialist with expertise in the diagnosis and management of XLH.

\(^1\)Normal range based on age and gender as informed by the Canadian Laboratory Initiative on Pediatric Reference Intervals (CALIPER; https://caliperproject.ca/) can be used as a reference of this information.

\(^2\)Baseline RSS total score is defined as the RSS total score measured immediately prior to initiation of Crysvita therapy.

Renewal Criteria:

Renewals of reimbursement will be considered in patients meeting the following criteria:

- Patient continues to be under the care of a specialist with expertise in the diagnosis and management of XLH; AND
- The patient does not meet any of the discontinuation criteria while on therapy.

Patients should be assessed on an annual basis. In pediatric patients in whom epiphyseal closure has not occurred, documentation demonstrating improvement from baseline RSS total score in the first 12 months and/or maintenance of RSS total score improvement achieved in the first 12 months is to be provided at the time of renewal.
Discontinuation Criteria:

- In pediatric\(^a\) patients in whom epiphyseal closure has not yet occurred and who met the above initiation criteria, treatment should be discontinued upon meeting any of the following circumstances:
  - There is no demonstrated improvement in the 12-month RSS total score from baseline\(^2\) RSS total score; OR
  - The patient’s RSS total score achieved after the first 12 months of burosumab therapy has not been maintained subsequently.

- In adolescent\(^b\) patients in whom epiphyseal closure has occurred and who met the above initiation criteria and initiated treatment as a pediatric patient, treatment should be discontinued upon development of any of the following clinical circumstances:
  - Hyperparathyroidism; OR
  - Nephrocalcinosis; OR
  - Evidence of fracture or pseudo-fracture based on radiographic assessment.

- In adult\(^c\) patients who met the above initiation criteria and initiated treatment as a pediatric patient, treatment should be discontinued upon development of any of the following clinical circumstances:
  - Hyperparathyroidism; OR
  - Nephrocalcinosis; OR
  - Evidence of fracture or pseudo-fracture based on radiographic assessment.

\(^2\)Baseline RSS total score is defined as the RSS total score measured immediately prior to initiation of Crysvita therapy.

\(^a\)Pediatric refers to a patient who is aged 17 years or younger.

\(^b\)Adolescent refers to a pediatric patient who is aged 13 to 17 years.

\(^c\)Adult refers to a patient who is 18 years of age and older.

**Approval duration for initial requests**: 1 year
**Burosumab**  
**Brand(s):** Crysvita  
**DOSAGE FORM/ STRENGTH:** 10 mg/mL, 20 mg/mL, 30 mg/mL vial

Approved dosage for initial and renewal:

Funding will be provided at a minimum dose of 10 mg up to a maximum dose of 90 mg.

- Children (1-17 years of age): 0.8mg/kg of body weight (rounded to the nearest 10 mg) subcutaneously every 2 weeks.
- Adults (18 years of age and older): 1mg/kg of body weight subcutaneously (rounded to the nearest 10 mg) every 4 weeks.

Dose to be adjusted based on serum phosphorus levels.
Canakinumab
Brand(s): Ilaris
DOSAGE FORM/ STRENGTH: 150 mg/vial injection

**Muckle-Wells Syndrome:**

The patient has a diagnosis of Muckle-Wells Syndrome (MWS) based on meeting each of the following confirmatory results:

1. Clinical diagnosis of MWS;
2. NLRP3 mutation (mutational analysis required);
3. SAA levels ≥ 10 mg/L; and
4. Assessment score of patient’s disease activity determined by the following 4 parameters: skin disease, arthralgia, conjunctivitis, and fatigue/malaise ≥ mild, moderate, or severe.

If a patient is without NLRP3 mutation, a request will be considered on a case-by-case basis if the other confirmatory criteria are met and if the exclusion criteria are not met. However, the initial approval would be for 6 months; if the patient demonstrated a sustained SAA level < 10 mg/L, a renewal of funding for a 1 year approval period would be considered.

Initial Approval Period: 1 year

**NOMID Syndrome:**

The patient has a Diagnosis of Neonatal-Onset Multisystem Inflammatory Disease (NOMID) based on meeting each of the following confirmatory results:

1. Clinical diagnosis of NOMID.
2. Presentation of symptoms of NOMID made in a patient < 6 months of age.
3. NLRP3 mutation (mutational analysis required).
4. A score of patient’s disease activity determined by the following 4 parameters: skin disease, arthralgia, conjunctivitis, and fatigue/malaise ≥ mild, moderate, or severe.

If a patient is without NLRP3 mutation, a request will be considered on a case-by-case basis if the other confirmatory criteria are met and if the exclusion criteria are not met. However, the initial approval would be for 6 months; if the patient demonstrated a sustained SAA level < 10 mg/L, a renewal of funding for a 1 year approval period would be considered.

Initial Approval Period: 6 months

**For Both Indications**

For all patients, the treatment is supervised by medical specialists with knowledge in the management of CAPS (Cryopyrin-Associated Periodic Syndromes).
Canakinumab
Brand(s): Ilaris
DOSAGE FORM/STRENGTH: 150 mg/vial injection

The approved dosage will be:

❖ Bodyweight > 40 kg: 150 mg subcutaneously every 8 weeks
❖ Bodyweight = 15 kg – 40 kg: 2 mg/kg subcutaneously every 8 weeks. (For children 15 kg – 40 kg with an inadequate response, the dose can be increased to 3 mg/kg.)

For all patients, funding requests will be denied if any of the exclusion criteria is met:

(i) The patient is bedridden where any physical activity brings on discomfort and symptoms which occur at rest AND not amenable to surgical/medical intervention;

(ii) The patient has another life-threatening disease where prognosis is unlikely to be influenced by Ilaris® (canakinumab) therapy (e.g. neuroblastoma, leukemia etc.);

(iii) The patient has a life-expectancy of six months or less, regardless of the cause; or

(iv) The patient has Familial Cold Auto-Inflammatory Syndrome (FCAS).

Renewal Criteria:

Confirmation from the Patient’s physician that the patient has benefited or continues to benefit from therapy with Ilaris, is expected to continue to do so and that each of the following is met:

1. The patient has SAA levels < 10 mg/L. If SAA levels 10 mg/L or higher over a 6 month interval and that is sustained over two consecutive 6 month intervals, approval for funding may be withdrawn.

2. An assessment score of patient’s skin disease, arthralgia, conjunctivitis, and fatigue/malaise shows no or minimal disease activity

3. The patient:
   (a) is not bedridden where any physical activity brings on discomfort and symptoms which occur at rest AND not amenable to surgical/medical intervention;
   (b) has no other life-threatening disease where prognosis is unlikely to be influenced by Ilaris® (canakinumab) therapy (e.g. neuroblastoma, leukemia etc.); or
Canakinumab
Brand(s): Ilaris
DOSAGE FORM/ STRENGTH: 150 mg/vial injection

(c) has not developed a life-threatening complication or a severe injection reaction to Ilaris® (canakinumab) not treatable by other therapeutic measures.

4. The patient has adhered with prescribed injection schedule for optimal management of the disease

5. The patient has adhered to all safety and effectiveness monitoring of the treatment

6. Treatment will be supervised by medical specialists with knowledge in the management of CAPS

The approved dosage for Subsequent Approvals is the same as Initial Approvals.
Cerliponase alfa
Brand(s): Brineura
DOSAGE FORM/ STRENGTH: 150mg/5mL (30mg/mL) Injection
Updated December 2, 2020

Initiation Criteria:
For the treatment of Neuronal Ceroid Lipofuscinosis Type 2 (CLN2) disease/ tripeptidyl peptidase 1 (TPP1) deficiency, in patients who meet the following criteria at the time of treatment initiation:

- Diagnosis of CLN2 disease is confirmed by TTP1 enzyme activity and CLN2 genotype analysis; AND
- CLN2 Rating Scale demonstrates the following requirements:
  - A minimum score greater than or the same as 1 in each of the motor and the language domains; AND
  - A minimum score greater than or the same as 3 for the aggregate motor-language score
- AND Patient is under the care of a prescriber/specialist with expertise in the diagnosis and management of CLN2 disease.

Approval duration of initials: 24 weeks

Renewal Criteria:

Renewal of funding will be considered for patients who do not meet any of the exclusion criteria and who have not demonstrated any of the stopping/discontinuation criteria while on therapy.

Exclusion Criteria (Applies to both initiation and renewal criteria):

Patients meeting any of the following criteria will not be funded:
- Requires ventilation support (except for non-invasive support at night);
- Presence of contraindications to intracerebroventricular (ICV) drug administration;
- Presence of acute ICV device complications;
- Presence of ventriculoperitoneal (VPS) shunts

Stopping/Discontinuation Criteria for Brineura:

Treatment with Brineura will be discontinued if:
- There is reduction of 2 or more points in the aggregate motor-language score of the CLN2 Clinical Rating Scale that is maintained over any 2 consecutive 24-week assessments; OR
- The aggregate motor-language score of the CLN2 Clinical Rating Scale reaches 0 (zero) at 2 consecutive 24-week assessments; OR
- The patient has developed contraindication to ICV drug administrations
Cerliponase alfa
Brand(s): Brineura
DOSAGE FORM/ STRENGTH: 150mg/5mL (30mg/mL) Injection

Recommended dose:

300 mg (10 mL solution) administered by intracerebroventricular (ICV) infusion once every 2 weeks.

Approval duration of renewals: 12 months
**Elosulfase Alfa**
**Brand(s):** Vimizim  
**DOSAGE FORM/ STRENGTH:** 1 mg/mL Injection

Initiation Criteria:

For the treatment of mucopolysaccharidosis type IVA (MPS IVA) in patients meeting all the following criteria;

i) 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 GALNS\(^1\); AND  
ii) Patient is under the care of a specialist with experience in the diagnosis and management of MPS IVA; AND  
iii) The following baseline evaluations prior to initiation of Vimizim (elosulfase alfa) must be provided with the request for coverage:  
   iv) Detailed medical history documenting surgeries, medical admissions, subspecialty assessments  
   v) Orthopedic evaluation including spinal and cranial MRI, skeletal x-rays, pain symptoms from bone and joints as appropriate to age and clinical disease.  
   vi) Mobility measure: 6MWT and stair climb (if appropriate for age and disease status)  
   vii) Respiratory function testing including sleep study testing (if appropriate for age)  
   viii) Age appropriate quality of life measure (such as HAQ, PODCI, EQ5D5L or SF36)\(^2\)  
   ix) documentation of mobility aide requirement, such as a walker or cane  
   x) documentation of requirement for respiratory aides, including ventilation status and changes in respiratory support requirements;  
   xi) Ophthalmologic and ear, nose and throat (ENT) assessment (if appropriate)  
   xii) Urine keratin sulfate (KS) determination: specific KS determination is preferred over total glycosaminoglycans (GAGs)  
   xiii) Cardiac echocardiogram

\(^1\)Note: not all MPS IVA patients will have 2 known pathogenic alleles identified and parental mutation analysis to establish the phase of mutations should be performed.

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

Exclusion Criteria (Patient will not be started on Vimizim if any of the following are met/apply):  

i) 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)  
ii) The patient has a forced vital capacity (FVC) of less than 0.3 liters and requires continuous ventilator assistance.  
iii) The patient/family is unwilling to comply with the associated monitoring criteria  
iv) The patient/family is unwilling to attend clinics for assessment and treatment purposes
Elosulfase alfa
Brand(s): Vimizim
DOSAGE FORM/ STRENGTH: 1 mg/mL

Approval duration of initials: 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
- Forced Vital Capacity (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 measure
- 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 continuation criteria
- Are unable to tolerate infusions due to infusion related adverse events that cannot be resolved
- Require permanent invasive ventilation
- Miss more than 6 infusions in a 12-month interval, unless for medically related issues.
- Meets any one of the Exclusion Criteria

Approval duration of renewals: 1 year

Recommended dose: 2mg/kg IV infusion once a week.
Glycerol phenylbutyrate
Brand(s): Ravicti
DOSAGE FORM/ STRENGTH: 1.1g/mL-25mL bottle

For the management of patients with chronic urea cycle disorders (UCD) who meet all the following criteria:

- Glycerol phenylbutyrate is being used as a nitrogen binding agent; AND
- Patient has demonstrated that they cannot be managed by dietary protein restriction and/or amino acid supplementation alone\(^1\); AND
- Patient is under the care of a physician with expertise in the treatment of patients with UCD or in consultation with a physician with this expertise.

\(^1\)The initial request should include levels for blood ammonia and glutamine levels demonstrating inadequate effects of protein restriction or amino acid supplementation.

Exclusion Criteria:

- Is not used in combination with other forms of phenylbutyrate
- Will not be funded for patients who are not using a low protein diet while on treatment
- Not funded for the management of acute hyperammonemia
- Not funded for patients under 2 months of age

Recommended dose: 5 g/m\(^2\) to 12.4 g/m\(^2\) per day

Approval duration: 1 year

Renewal Criteria:

Renewals will be considered in patients who demonstrate benefit from treatment\(^2\) and who have not developed unacceptable toxicities requiring discontinuation.

\(^2\)At the time of renewal, please provide recent (within 3 months) blood ammonia and glutamine levels while on treatment and address the number and severity of hyperammonemic events experienced while on treatment in the previous 12 months and any treatment emergent events requiring urgent care or hospitalization.

First renewal: 1 year

Subsequent renewals: 2 years
Initiation criteria

For the treatment of polyneuropathy in patients with hereditary transthyretin-mediated amyloidosis (hATTR), meeting all the following criteria:

1. Age 18 years of age or older; AND
2. Has a confirmed genetic diagnosis of hereditary transthyretin-mediated amyloidosis: AND
3. Symptomatic with Polyneuropathy disability stage I to ≤ IIIB or with Familial amyloidotic polyneuropathy stage I or II.
4. Under the care of a specialist with experience in the diagnosis and management of hATTR

Exclusion Criteria:
- Pre-symptomatic patients
- Patients diagnosed with severe heart failure symptoms (defined as New York Heart Association class III or IV).
- Patients who are recipients of a liver transplant
- Patients with platelet count < 100 x 10^9/L before initiation of treatment
- Patients who will be using Inotersen in combination with other interfering ribonucleic acid drugs or transthyretin stabilizers used to treat hATTR.

Discontinuation criteria:

Treatment with inotersen will be discontinued for patients who are:
- Permanently bedridden and dependent on assistance for basic activities of daily living, or
- Receiving end-of-life/palliative care where survival of less than one year is expected.

Renewal Criteria:

Renewal of funding will be considered if patients do not meet the discontinuation criteria.

Dosage: 300 mg of inotersen sodium (284mg of inotersen) sc injection once a week.

Patients should be assessed after 9 months of treatment and then every six months thereafter.

Duration of Approval of initiation requests: 10 months
Duration of Approval of first renewal: 6 months
Duration of Approval of 2nd and subsequent renewals: 1 year
Inotersen
Brand(s): Tegsedi
DOSAGE FORM/ STRENGTH: 284 mg/ 1.5 mL prefilled syringes

Notes to Prescribers:
- Laboratory documentation for the genetic mutation for hATTR must be included with the application.
- Signs and symptoms of polyneuropathy should be listed.
- In your application please list all drugs that the patient is using including whether he/she is using any of the following: Diflunisal, Patisiran, Tafamidis
- Confirmation that the patient does not meet each of the listed exclusions must be provided on the request.

Definitions:

Familial Amyloid Polyneuropathy (FAP) stage: Clinical staging system for the neuropathy symptoms of hATTR (formerly termed familial amyloid neuropathy).
- FAP Stage 1: Walking without assistance, mild neuropathy (sensory, autonomic, and motor) in lower limbs
- FAP Stage 2: Walking with assistance, moderate impairment in lower limbs, trunk, and upper limbs
- FAP Stage 3: wheelchair or bed-ridden, severe neuropathy

Polyneuropathy disability score (PND): A five-stage measure of neuropathy impairment ranging from 0 (no impairment) to 4 (confined to a wheelchair or bedridden).
- Stage 0: no impairment
- Stage I: sensory disturbances but preserved walking capability
- Stage II: impaired walking capability but ability to walk without a stick or crutches
- Stage IIIA: walking only with the help of one stick or crutch
- Stage IIIB: walking with the help of two sticks or crutches
- Stage IV: confined to a wheelchair or bedridden
EAP Initiation criteria:

For the treatment of Fabry disease (FD) in adult patients who meet ALL the following criteria;
1. Has a confirmed diagnosis of Fabry disease (deficiency of α-galactosidase A [α-Gal A]) and must be otherwise eligible for enzyme replacement therapy (ERT) for the treatment of FD as determined by a panel of Fabry disease experts consistent with criteria for diagnosis from the Canadian Fabry Disease Initiative (CFDI)\(^1\)
2. Has an α-galactosidase A mutation that is determined to be amenable to migalastat by an in vitro assay\(^2\)
3. Migalastat is not being used concomitantly with ERT\(^3\).
4. Patient is considered to be compliant/adherent to treatment
5. Prescriber must be an expert in genetic disorders or a clinician experienced in the diagnosis and management of Fabry disease.

\(^1\)Refer to Canadian Fabry Disease treatment guidelines 2017 at

Information submitted with requests to support the diagnosis of FD should include:

i) Clinical features associated with FD
ii) Biochemical markers (e.g. alpha-galactosidase activity in plasma or leukocytes, elevated plasma and/or urine biomarkers)
iii) Molecular changes
iv) Pathologic findings (e.g. biopsy results from involved tissues)

\(^2\)Definition of an amenable mutation: mutation that increases activity of alpha galactosidase A in an in vitro cell culture system (human embryonic kidney or HEK cells) by 1.2 times the baseline activity with an absolute value for enzyme activity of 3% or greater when compared with wildtype values. You may refer to the following website http://canada.galafoldamenabilitytable.com/?validated=1&redirect=en&hpc=1&licensed=1 or other appropriate supplementary tables to determine amenable α-Gal A mutations to migalastat.

\(^3\)Enzyme replacement therapies (ERT) for Fabry disease (e.g. agalsidase alfa [Replagal, agalsidase beta [Fabrazyme])

Exclusion criteria:

- Individuals who do not have an amenable mutation to migalastat
- If used concomitantly with an enzyme replacement therapy for FD
- Individuals with severe renal insufficiency (GFR <30 mL/min/1.73m\(^2\))
- Individuals who are pregnant or nursing
- Individuals with poor adherence/compliance to treatment
Migalastat
Brand(s): Galafold
DOSAGE FORM/ STRENGTH: 123 mg capsule

**Note:** Patients should be monitored every 6 months or more frequently during the first 3 to 5 years of treatment.

**Recommended dose:** 123 mg every other day

**Approval duration:** 6 months

**Renewal criteria:**

Renewals will be considered on a case-by-case basis in individuals who;
- are adherent with treatment; and
- who demonstrate a response while on therapy as compared to baseline results; and
- who have not developed unacceptable toxicities to migalastat; and
- who continue not meet any of the exclusion criteria (see above)

Requests for ongoing reimbursement should include information related to the patient’s renal and cardiac function, cerebrovascular events, hospitalizations or emergency room visits for FD-related issues, health related quality of life measures as evaluated through a valid HRQL test (SF-36), gastrointestinal symptoms, pain measures, and other relevant clinical outcomes from treatment. Please address the patient’s adherence/compliance with treatment and any adverse effects / toxicities from treatment with migalastat. The Fabry disease expert panel may be consulted for renewal recommendations.

**Approval duration of renewals:** 1 year
For the treatment of spinal muscular atrophy (SMA) in patients meeting all the following criteria:

a) Provides genetic documentation of 5q SMA homozygous gene deletion, homozygous mutation, or compound heterozygote.

b) Patient meets one (1) of the following clinical scenarios:
   - Pre-symptomatic with 2 or 3 copies of the survival motor neuron 2 (SMN2) gene; OR
   - Have had disease duration of less than 6 months, 2 copies of the SMN2 gene, and symptom onset after the first week after birth and on or before 7 months of age; OR
   - Are under the age of 18 with symptom onset after 6 months of age, and never achieved the ability to walk independently.

c) Patient does not require permanent invasive ventilation.

d) Patient is under the care of a specialist experienced in the diagnosis and management of SMA.

In addition, symptomatic Type 2 and 3 patients under the age of 18 regardless of ever achieving the ability to walk independently will be considered on a case by case basis.

Other patients who do not meet the expanded funding criteria may be considered in exceptional cases.

Renewal Criteria:
Renewal of funding will be considered for patients who have not demonstrated any of the Stopping/discontinuation criteria while on therapy.

Stopping/Discontinuation Criteria for Spinraza

These criteria are applicable to patients funded upon meeting either initial or renewal criteria.

An assessment of the response to therapy should be made prior to the fifth dose or every subsequent dose of Spinraza. Treatment should be discontinued upon meeting any of the following circumstances:

- Patient is not demonstrating/achieving response in motor milestones as assessed using 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), as follows:
  - there is no demonstrated improvement in motor milestone function above pre-treatment scores (as assessed using the HINE Section 2, CHOP INTEND or HFMSE); OR
Nusinersen
Brand(s): Spinraza
DOSAGE FORM/ STRENGTH: 2.4mg/mL – 5mL Vial Pk

- there is no demonstrated maintenance of motor milestone function (as assessed using HINE Section 2, CHOP INTEND or HFMSE);

OR

- Patient requires Permanent Invasive Ventilation (PIV)\(^1\)

Exclusion criteria:

a) Patient has SMA type 4
b) Patient has more than four (4) SMN2 gene copies
c) Patient with permanent invasive ventilation (PIV)\(^1\)

\(^1\)Permanent Invasive Ventilation (PIV) 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.

Recommended dose:

Loading doses:

12 mg administered intrathecally on days 0, 14, 28 and 58 (Note: 4\(^{th}\) loading dose is administered approximately 30 days after the 3\(^{rd}\) loading dose)

Maintenance dose:

12 mg administered intrathecally every 4 months starting 4 months after the 4\(^{th}\) loading dose.

Approval duration for initial request: 8 months

Approval duration of Renewals: 1 year
Onasemnogene abeparvovec
Brand(s): Zolgensma
DOSAGE FORM/ STRENGTH: 2 x 10^{13} vector genomes (VG)/mL solution for infusion
Effective date: October 29, 2021

For the treatment of spinal muscular atrophy (SMA) in individuals meeting all the following criteria:
1. Diagnosis of SMA is confirmed by genetic documentation of 5q spinal muscular atrophy with biallelic mutations in the survival motor neuron 1 (SMN1) gene; AND
2. Patient is 180 days of age or younger at the time that the treatment is administered; AND
3. Patient is pre-symptomatic or symptomatic with one to three copies of the survival motor neuron 2 (SMN2) gene; AND
4. Patient does not require permanent ventilatory support (invasive or non-invasive)* or a permanent feeding tube; AND
5. Patient must be under the care of a specialist with experience in the diagnosis and management of spinal muscular atrophy.

*Permanent ventilatory support is defined as the need for a tracheostomy or requirement of 16 hours or more of respiratory assistance per day (via non-invasive ventilatory support) for 14 or more consecutive days in the absence of an acute reversible illness excluding perioperative ventilation.

Exclusion Criteria:
1. Reimbursement is limited to one lifetime administration of onasemnogene abeparvovec. Patients who have received a prior dose of onasemnogene abeparvovec accessed by any mechanism (e.g. private insurance plan, clinical trial, compassionate access) will not be funded.
2. Patients with 4 (four) or more copies of SMN2 gene will not be funded.

Notes:
1. Patients must test for the presence of anti-AAV9 antibodies and provide the report of antibody titers. It should be noted that patients with titers above 1:50 will not be funded. Repeat testing results will be considered for newborns suspected of having acquired interim antibodies from maternal transmission.
2. No further treatment with nusinersen or other medications indicated for the treatment of SMA will be considered after the patient has received a dose of onasemnogene abeparvovec.

Dosage recommended: A maximum single lifetime dose of 1.1 x 10^{14} vector genomes (vg)/kg
Renewal of therapy is not considered.
Patisiran
Brand(s): Onpattro
DOSAGE FORM/ STRENGTH: 2 mg/mL injection
Effective date: April 8, 2021

Initiation criteria

For the treatment of polyneuropathy in patients with hereditary transthyretin-mediated amyloidosis (hATTR), meeting all the following criteria:

1. Age 18 years of age or older; AND
2. Has a confirmed genetic diagnosis of hereditary transthyretin-mediated amyloidosis: AND
3. Symptomatic with Polyneuropathy disability stage I to ≤ IIIB or with Familial amyloidotic polyneuropathy stage I or II.
4. Under the care of a specialist with experience in the diagnosis and management of hATTR

Exclusion Criteria:
• Pre-symptomatic patients
• Patients diagnosed with severe heart failure symptoms (defined as New York Heart Association class III or IV).
• Patients who are recipients of a liver transplant
• Patients who will be using patisiran in combination with other interfering ribonucleic acid drugs or transthyretin stabilizers used to treat hATTR.

Discontinuation criteria:

Treatment with patisiran will be discontinued for patients who are:
• Permanently bedridden and dependent on assistance for basic activities of daily living, or
• Receiving end-of-life/palliative care where survival of less than one year is expected.

Renewal Criteria:
Renewal of funding will be considered if patients do not meet the discontinuation criteria.

Dosage: 0.3 mg/kg IV once every three weeks, with a maximum dose of 30 mg for patients who weigh 100 kg or greater.

Patients should be assessed after 9 months of treatment and then every six months thereafter.

Duration of Approval of initiation requests: 10 months
Duration of Approval of first renewal: 6 months
Duration of Approval of 2nd and subsequent renewals: 1 year

Notes to Prescribers:
• Laboratory documentation for the genetic mutation for hATTR must be included with the application.
• Signs and symptoms of polyneuropathy should be listed.
• In your application please list all drugs that the patient is using including whether he/she is using any of the following: Diflunisal, Inotersen, Tafamidis
• Confirmation that the patient does not meet each of the listed exclusions must be provided on the request.

Patisiran
Brand(s): Onpattro
DOSAGE FORM/ STRENGTH: 2 mg/mL Injection

Definitions:

Familial Amyloid Polyneuropathy (FAP) stage: Clinical staging system for the neuropathy symptoms of hATTR (formerly termed familial amyloid neuropathy).
- FAP Stage 1: Walking without assistance, mild neuropathy (sensory, autonomic, and motor) in lower limbs
- FAP Stage 2: Walking with assistance, moderate impairment in lower limbs, trunk, and upper limbs
- FAP Stage 3: wheelchair or bedridden, severe neuropathy

Polyneuropathy disability score (PND): A five-stage measure of neuropathy impairment ranging from 0 (no impairment) to 4 (confined to a wheelchair or bedridden).
- Stage 0: no impairment
- Stage I: sensory disturbances but preserved walking capability
- Stage II: impaired walking capability but ability to walk without a stick or crutches
- Stage IIIA: walking only with the help of one stick or crutch
- Stage IIIB: walking with the help of two sticks or crutches
- Stage IV: confined to a wheelchair or bedridden
Initiation Criteria:

For the treatment of patients with proven acromegaly who meet the following criteria;

1. Active disease as indicated by GH concentration following an oral glucose tolerance test of >1 ug/L; AND

2. Failed pituitary surgery or pituitary surgery is not possible\(^1\); AND

3. After a failed 6 month trial of a somatostatin analogue\(^2\) (Failure is defined as IGF-1 levels more than 25% above upper limits of age-adjusted normal range.)

4. Treatment should be supervised by an endocrinologist.

Note: Maximum daily dose of 30mg of Pegvisomant will be approved.

Approval Duration: 1 year

Renewal Criteria:

Patient has been able to tolerate the medication (i.e. no significant adverse effects) and there is objective evidence of response to therapy demonstrated by:

1. Normalization of IGF-1 level; AND

2. Improvement in the patient’s symptoms and/or co-morbid complications; AND

3. No progression of pituitary tumour

Note: Maximum daily dose of 30mg of Pegvisomant will be approved. The approval letter will include a note to consider once weekly dosing.

Approval Duration: 1 year

\(^1\)Surgery may not be appropriate in some patients due to technical reasons or due to unstable co-morbid conditions. The requesting physician should provide documentation (i.e. surgical consultation notes). Patients with acromegaly due to non-pituitary tumours will also be considered for reimbursement using the above criteria.

\(^2\)Patient must have documented intolerance to the maximal dose and/or have failed to achieve normalization of age-adjusted IGF-1 levels from treatment with maximal dose.
Ongoing funding of sapropterin (Kuvan) will be considered through the EAP for non-pregnant 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 [generally reimbursed through the Biomarin, the manufacturer of Kuvan].

**Initial Criteria for the Trial Period (First 6 months)**

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:

1. Diagnosis of PKU confirmed through an approved test.
2. Compliance with low protein diet and formulas.
3. Documented baseline blood phenylalanine (Phe) levels, which have been measured at least twice during a 3 to 6 month period, are greater than 360 μmol/L despite compliance with a low protein diet;
4. Baseline protein intake assessment by a dietitian. Ability to comply with their medication regimen.
5. Managed by a prescriber specialized in metabolic/biochemical diseases.
6. The Patient’s blood Phe tolerance levels will be documented at months 1 to 2 and 4 to 6 during the Trial Period.

**Modified Criteria for Pregnant Patients** during the 6 month trial period:

1. Patient has a diagnosis of PKU confirmed through an approved test.
2. Patient’s treatment is being managed by a prescriber specialized in metabolic/biochemical diseases; and
3. 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.

In the case of Patients who are eligible for but do not utilize the Patient support program for a 185-day or 6-month trial. Executive Officer will approve a request to reimburse claims for Kuvan at a dosage of up to 20mg/kg per day for the trial period of up to 6 months provided that the above conditions are met.
Sapropterin
Brand(s): Kuvan
DOSAGE FORM/ STRENGTH: 100 mg tablet

Approval Duration: 6 months

Funding Criteria for Kuvan in the Post Trial Period: Patients must have demonstrated the response as per the trial criteria to be funded following the 6 month trial period.

Initial Criteria Post 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:

1. Compliance with low protein diet, formulas, and Kuvan; AND

2. During the 6 month trial period under EAP or PSP patient has achieved one of more of the following:
   i) a demonstrated response to the Kuvan responsiveness test or PKU clinical protocol, based on the following information:
   ii) the clinic’s definition for response; and
   iii) all relevant laboratory results used to determine that the Patient was a responder to Kuvan

3. Patient meets one of the following:
   i) normal sustained Blood Phe levels [ > 120 μmol/L and < 360 μmol/L] (At least 2 levels measured at least 1 month apart); OR
   ii) 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
   iii) 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;

4. Demonstrated an increase in dietary protein tolerance based on targets set between the clinician and patient

5. Managed by a prescriber specialized in metabolic/ biochemical diseases.

Dosage: Up to a maximum of 20 mg/kg per day

Approval Duration: 1 year
Sapropterin  
Brand(s): Kuvan  
DOSAGE FORM/ STRENGTH: 100 mg tablet

Renewal Criteria:
Renewals will be considered for patients meeting the following criteria:
1. Demonstrates ongoing response to treatment; AND
2. Complies with and will continue to comply with a low protein diet, formulas, and treatment with Kuvan; AND
3. the request for extended coverage includes a recent follow-up from a prescriber specialized in metabolic/biochemical diseases.

Exclusion Criteria for both Initial (Trial and Post Trial period) and Renewal criteria: (Patients meeting any of the following criteria will not be funded)
- the Patient is not on a low protein diet or is not compliant with their low protein diet; or
- the Patient has a baseline blood PHe level less than 360 μmol/L prior to initiating therapy with Kuvan.

Dosage: Up to a maximum of 20 mg/kg per day

Approval Duration: 1 year
Sebelipase alfa
Brand(s): Kanuma
DOSAGE FORM/ STRENGTH: 2 mg/mL solution
Effective date: April 8, 2021

Initiation Criteria:

For the treatment of lysosomal acid lipase (LAL) deficiency in patients meeting ALL the following criteria:

- Has documented biochemical evidence of deficient LAL activity; AND
- Has two documented pathogenic mutations in the LIPA gene; AND
- Is under the care of a specialist with experience in the diagnosis and management of LAL deficiency; AND
- The onset of clinical manifestations of LAL deficiency occurred before six months of age;

OR

Patient is six months of age or older; AND

- Presents with one or more of the following:
  - Persistently elevated transaminases (Alanine aminotransferase (ALT) > 1.5 x ULN\(^1\) or Aspartate transaminase (AST) > 1.5 x ULN\(^1\)) as measured by two assessments three to six months apart
  - Persistent dyslipidemia (Low-density lipoprotein cholesterol (LDL-C) and/or Triglycerides (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 impairment\(^2\)
  - Evidence of intestinal affection and/or malabsorption;

AND

- Does not present with 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); OR
  - Severe hepatic dysfunction (Child-Pugh Class C); OR
  - End-stage liver disease.

NOTE: The requesting prescriber must provide baseline values for the following clinical components before starting sebelipase alfa; Bloodwork for ALT, AST, liver fibrosis, growth curve, spleen and liver volume/sizes, portal vein pressures, patient’s age, weight, and height.
Discontinuation Criteria:

Reimbursement will be discontinued in patients who have experienced adverse events from sebelipase alfa (particularly hypersensitivity reactions, including anaphylaxis, hypotension, or fever), which cannot be managed with standard treatment, and/or which have a significant impact on the patient’s quality of life, or are life-threatening.

For patients with onset of clinical manifestations of LAL deficiency at six months of age and older, reimbursement will be discontinued if:

- Patient has progressed to end-stage liver failure or multi-organ failure; OR
- Patient has at least three out of the five following response components compared with baseline values after 12 months of therapy:
  1. Less than 10% improvement in ALT or AST
  2. Worsening of liver fibrosis confirmed by biopsy
  3. Persisting growth impairment\(^2\) despite sebelipase alfa therapy and nutritional interventions
  4. At least a 15% increase in spleen volume and/or greater than 15% increase in liver volume on ultrasound
  5. Increased portal vein pressures, or de novo evidence of portal hypertension on ultrasound and Doppler, or new presentation of portal hypertension (e.g. esophageal varices)

\(^1\) Based on age- and sex-specific normal values for ALT and AST

\(^2\) Growth impairment is defined as decreased body weight across at least two of the major centiles on a World Health Organization (WHO) weight-for-age chart, or body weight below 10\(^{th}\) 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 (URLs: [https://www.who.int/toolkits/child-growth-standards/standards/weight-for-age](https://www.who.int/toolkits/child-growth-standards/standards/weight-for-age), [https://www.who.int/toolkits/child-growth-standards/standards/length-height-for-age](https://www.who.int/toolkits/child-growth-standards/standards/length-height-for-age))

Exclusion Criteria: Patients with severe liver disease and/or those who have progressed to end stage liver disease.

Renewal Criteria: Renewals will be considered in patients who do not meet the discontinuation criteria.

Recommended dose:

For patients with onset of clinical manifestations of LAL deficiency before six months of age:

1mg/kg administered as an intravenous infusion once weekly. Based on clinical response, dose increase to 3mg/kg may be considered.
For patients with onset of clinical manifestations of LAL deficiency at six months of age or older:

1mg/kg administered as an intravenous infusion once every other week.

Approval duration of initials and renewals: 12 months initial, 6 months renewal
Initiation criteria

For the treatment of adult patients with cardiomyopathy due to transthyretin (TTR)-mediated amyloidosis in patients meeting the following criteria:

1. Patient has documented cardiac disease due to wild-type or hereditary TTR-mediated amyloidosis cardiomyopathy (ATTR-CM)
   i) Documented wild-type ATTR-CM consists of all of the following:
      • absence of a variant TTR genotype;
      • evidence of cardiac involvement by echocardiography with end-diastolic interventricular septal wall thickness >12 mm;
      • presence of amyloid deposits in biopsy tissue (fat aspirate, salivary gland, median nerve connection tissue sheath, or cardiac); and
      • TTR precursor protein identification by immunohistochemistry, scintigraphy, or mass spectrometry.
   ii) Documented hereditary ATTR-CM consists of all of the following:
      • presence of a variant TTR genotype associated with cardiomyopathy and presenting with a cardiomyopathy phenotype;
      • evidence of cardiac involvement by echocardiography with end-diastolic interventricular septal wall thickness greater than12 mm;
      • presence of amyloid deposits in biopsy tissue (fat aspirate, salivary gland, median nerve connective tissue sheath, or cardiac)

2. Patient with New York heart Association (NYHA) class I to III

3. History of heart failure, defined as at least one prior hospitalization for heart failure or clinical evidence of heart failure that required treatment with a diuretic.

4. Request is from a prescriber with experience in the diagnosis and management of ATTR-CM.

Exclusion criteria:

Patients meeting one or more of the following will not be eligible for funding:
   • Patients who have received a heart or liver transplant
   • Patients with an implanted cardiac mechanical assist device (CMAD)
   • Patients being treated with other disease-modifying treatments for ATTR
   • Patients with NYHA Class IV
Tafamidis
Brand(s): Vyndaqel
DOSAGE FORM/ STRENGTH: 20 mg capsule

Discontinuation criteria:

Treatment with tafamidis will not be funded upon meeting one or more of the following discontinuation criteria:

• Patients who progress to NYHA Class IV
• Patients who receive a heart or liver transplant
• Patient who receive an implanted CMAD

Renewal criteria: Renewals will be considered in patients who do not meet the discontinuation criteria.

Recommended dose: 80 mg (administered as four 20 mg capsules) taken orally, once daily, with or without food. Dose may be adjusted to a dose of 20mg if not tolerated.

Initial approval duration: 6 months
First renewal duration: 6 months
Second and subsequent renewal duration: 1 year
MIGRAINE DRUGS

Onabotulinum Toxin A
Brand(s): Botox
DOSAGE FORM/ STRENGTH: 50 U/Vial, 100 U/Vial, 200 U/vial

For the prophylaxis of headaches in adults meeting the following criteria for funding:

- Patient with chronic migraine (defined as ≥15 days per month with continuous headache lasting ≥4 hours AND at least 4 distinct headache episodes each lasting ≥4 hours); AND

- Patient has failed¹ three or more prior oral prophylactic medications²; AND

- Request for Botox to treat migraine must be provided by a physician with specialty training in the management of headache. Administration should only be given by physicians with the appropriate qualifications and experience in the treatment, use, and proper administration of Botox for headaches.

¹Failure is defined as no therapeutic or unsatisfactory effect (Less than a 30% reduction in frequency of headache days) to an adequate dose and duration of 3 prophylactic therapies² where two treatments must be of different types/classes.

Contraindication or intolerable side effects necessitating discontinuation will be considered for 1 of the 3 drugs only.

²Prophylactic therapies to be considered include:

- Beta blockers
- Tricyclic antidepressants
- Verapamil or flunarizine
- Sodium valproate (or divalproex sodium)
- Topiramate
- Gabapentin

Requests should contain the following information:

- Objective measure of baseline headache days and response to other prophylactic medications (i.e. headache diary)
- List of previously tried prophylactic medications, including doses and duration as well as why they were discontinued
- Confirmation of specialty training in the management of headache.
Onabotulinum Toxin A  
Brand(s): Botox  
DOSAGE FORM/ STRENGTH: 50 U/Vial, 100 U/Vial, 200 U/vial  

Dosing: As per product monograph  
Notes regarding continued therapy with “Botox”:  

i) Patients who have not obtained an adequate treatment response after 2 treatment cycles should be discontinued from further therapy.  

ii) Patients who obtain an adequate response and who transition from chronic migraine to episodic migraine should be discontinued from therapy within 3 months of that transition.  

An adequate treatment response is defined as a ≥ 50% reduction in frequency of headache days per month  

Duration of Approval: 1 year  

Renewal criteria:  

• Objective evidence (i.e. headache diary) that the patient has obtained an adequate treatment response defined as a ≥ 50% reduction in frequency of headache days per month; AND  

• Confirmation that the patient has not transitioned from chronic migraine to episodic migraine. Therapy will be reimbursed for a maximum of 3 months after transition from chronic migraine to episodic migraine.  

• Consideration will be given for renewals in patients who had an initial adequate response to Botox, discontinued therapy and subsequently transitioned back to chronic migraine status.  

Duration of Approval: 1 year
### Almotriptan
**Brand(s):** Axert  
**DOSAGE FORM/ STRENGTH:** 6 mg, 12.5 mg tablet

### Naratriptan
**Brand(s):** Amerge  
**DOSAGE FORM/ STRENGTH:** 1 mg, 2.5 mg tablet

### Rizatriptan
**Brand(s):** Maxalt, Maxalt RPD  
**DOSAGE FORM/ STRENGTH:** 5 mg, 10 mg tablet and wafer

### Sumatriptan
**Brand(s):** Imitrex  
**DOSAGE FORM/ STRENGTH:** 50 mg, 100 mg tablet

For the treatment of migraines with or without aura in patients who failed adequate trials of other medications for migraines (e.g. acetaminophen, NSAIDs) and where the following information is provided:

- Details of migraine prophylactic regimens (e.g. amitriptyline, beta-blockers) tried or rationale why they are inappropriate; and
- The number of attacks, duration, and severity of migraines.

**Duration of Approval:** 5 years

**Renewal** requests may be considered for patients who continue to benefit from treatment. The physician must provide the frequency of triptan use.

*Warning: The frequent use of triptans (i.e. more than three days per week for longer than three months at a time) may predispose a patient to developing triptan-induced chronic daily headaches.*
**Sumatriptan**

Brand(s): Imitrex Injection  
**DOSAGE FORM/ STRENGTH:** 12 mg/mL subcutaneous injection

Brand(s): Imitrex Nasal Spray  
**DOSAGE FORM/ STRENGTH:** 5 mg/dose and 20 mg/dose nasal spray

For the treatment of migraines with or without aura in patients who failed adequate trials of other medications for migraines (e.g. acetaminophen, NSAIDs) and has documented intolerance* to an oral triptan. The following information must also be provided:

- Details of migraine prophylactic regimens (e.g. amitriptyline, beta-blockers) tried or rationale why they are inappropriate; and
- The number of attacks, duration, and severity of migraines.
- The nature of intolerance or why oral sumatriptan cannot be used must be specified.

Duration of Approval 5 years

**Renewal** requests for sumatriptan may be considered for patients who continue to benefit from treatment. The physician must provide the frequency of triptan use.

*Warning: The frequent use of triptans (i.e. more than three days per week for longer than three months at a time) may predispose a patient to developing triptan-induced chronic daily headaches.*

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**Zolmitriptan**

Brand(s): Zomig  
**DOSAGE FORM/ STRENGTH:** 2.5 mg tablet

Brand(s): Zomig Rapimelt  
**DOSAGE FORM/ STRENGTH:** 2.5 mg dispersible tablet

For the treatment of migraines with or without aura in patients who have failed an adequate trial of or experienced intolerance to all other oral triptans considered under the Exceptional Access Program.

Duration of Approvals: 5 years

**Renewal** requests may be considered for patients who continue to benefit from treatment. The physician must provide the frequency of triptan use.
MULTIPLE SCLEROSIS DRUGS

Alemtuzumab
Brand(s): Lemtrada
DOSAGE FORM/ STRENGTH: 12 mg/ 1.2 mL Solution for IV infusion

For the treatment of Relapsing–Remitting Multiple Sclerosis (RRMS) as monotherapy in patients who meet all of the following criteria:

- The patient’s physician provides documentation setting out the details of the patient’s most recent neurological examination (which must have been conducted within ninety [90] days of the request, including a description of any recent attacks, the dates of attacks, and neurological findings); AND

- Patient has failed to respond¹ to full and adequate courses of at least one of the following therapies: interferon, glatiramer acetate, dimethyl fumarate, or teriflunomide or ocrelizumab has had a documented intolerance or contraindication to TWO or more of the listed therapies; AND

- Patient has experienced one (1) or more clinically disabling relapses in the previous year; AND

- Patient has had a 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;

- Patient is being followed by a neurologist experienced in the management of relapsing–remitting multiple sclerosis (RRMS); AND

- The patient has a current Expanded Disability Status Scale (EDSS) score less than or equal to 5.0.

¹ “Failed to respond to full and adequate courses” of certain therapies means that the patient has received a trial of at least 6 months of interferon, glatiramer acetate, dimethyl fumarate therapy, or teriflunomide; AND has experienced at least one disabling relapse (attack) while on such therapy.

Exclusion Criteria: No reimbursement if the patient satisfies any of the following exclusion criteria:

- the patient is receiving combination therapy of Lemtrada with other disease modifying therapies, such as Aubagio, Avonex, Betaseron, Copaxone/Glatect, Extavia, Rebif, Extavia, Tysabri, Gilenya, Ocrevus and Tecfidera; OR

- the patient has an EDSS score greater than 5.0; OR

- the patient is younger than 18 years old.

Dosage: 12 mg per day for two treatment courses.
**Alemtuzumab**  
**Brand(s):** Lemtrada  
**DOSAGE FORM/ STRENGTH:** 12 mg/ 1.2 mL Solution for IV infusion

Initial course: 12 mg per day for 5 consecutive days (60mg total dose). Second course: 12 mg per day for 3 consecutive days (36mg total dose) administered 12 months after the initial treatment course.

Retreatment beyond two cycles (eight vials) may be considered.  
**Note:** MRI reports are NOT mandatory to submit with the initial request.

**Cladribine**  
**Brand(s):** Mavenclad  
**DOSAGE FORM/ STRENGTH:** 10 mg tablet

**Initiation Criteria**

For the treatment of Relapsing Remitting Multiple Sclerosis (RRMS) in adult patients with active disease meeting ALL the following criteria:

1. 18 years of age or older
2. Diagnosis of RRMS is in accordance with the McDonald 2017 criteria demonstrating dissemination of lesions in the central nervous system in space and time meeting the following:
   i) 2 or more attacks\(^1\) and clinical evidence of 2 or more lesions\(^2\);  
      OR
   ii) 2 or more attacks\(^1\) and clinical evidence of 1 lesion with clear historical evidence of prior attack involving lesion in different location;

\(^1\)If the patient has experienced only one attack, the patient must meet ONE of the additional criteria of dissemination in time in the list below:
   • Additional clinical attack
   • Simultaneous presence of both enhancing and non-enhancing, symptomatic or asymptomatic MS-typical MRI lesions; OR new T2 or enhancing MRI lesion compared to baseline scan (without regard to timing of baseline scan)
   • Presence of cerebrospinal fluid (CSF)-specific oligoclonal bands

\(^2\)If the patient has evidence of only one lesion the patient must meet ONE of the additional criteria of dissemination in space in the list below:
   • Additional clinical attack implicating different CNS site
   • 1 or more MS-typical T2 lesions in 2 or more areas of the Central Nervous System (CNS): periventricular, cortical, juxtacortical, infratentorial or spinal cord  
   AND
Cladribine  
Brand(s): Mavenclad  
DOSAGE FORM/STRENGTH: 10 mg tablet

3. Failure or documented intolerance to at least one of an interferon OR glatiramer acetate OR dimethyl fumarate OR teriflunomide OR ocrelizumab; AND

4. Patient has experienced a clinical relapse and/or new MS lesions in the last 2 years; AND

5. Patient has an EDSS score less than 6.0 before start of therapy; AND

6. Cladribine is used as monotherapy for the treatment of RRMS; AND

7. The drug request is from a neurologist experienced in the management of RRMS or includes a consult note from a neurologist from an MS clinical recognized by the MS Society of Canada supporting the diagnosis.

Requests from community neurologists not working with an MS Clinic will be considered case-by-case. (Please include MRI reports or labwork to support the diagnosis of RRMS aligned with McDonald 2017 criteria.)

*Note: Requests for patients who is under the care of a community neurologist working outside of one of the MS Society recognized Ontario MS clinics can be considered on a case-by-case basis. Submit MRI and relevant history with your request.

Exclusion criteria:

1. Combination therapy with another disease modifying therapy for RRMS will not be reimbursed.

2. Patients with an EDSS score equal to or greater than 7.0

Dosage: Refer to the Mavenclad product monograph for dosing regimen based on body weight.

Taken as two treatment courses over 2 years. Each treatment course consists of 2 treatment weeks, which are one month apart at the beginning of each treatment year. 1.75 mg/kg per year administered as a treatment course in year 1 and a second treatment course starting 12 months after the first course in the respective year upon monitoring for recovery of lymphocytes before the second course of treatment is administered in accordance with prescribing information.

The recommended cumulative dose of is 3.5 mg/kg body weight over 2 years (1.75 mg/kg per year).

Approval Duration: Two treatment courses over 2 years are funded
For the treatment of Relapsing–Remitting Multiple Sclerosis (RRMS) in patients who meet all of the following criteria:

- The patient’s physician provides documentation setting out the details of the patient’s most recent neurological examination (which must have been conducted within ninety [90] days of the request, including a description of any recent attacks, the dates of attacks, and neurological findings).
- Patient has had one (1) or more clinical relapses in the previous year.
- The drug is requested by and followed by a neurologist experienced in the management of RRMS.
- The patient has a recent Expanded Disability Status Scale (EDSS) score ≤ 5.

Dosage: Initial: 120 mg twice daily
Maintenance: 240 mg twice daily

Renewal requests will be considered. Renewals for Tecfidera can be submitted through the Telephone Request Service.

The date and details of the most recent neurological examination and EDSS scores must be provided (exam must have occurred within the last ninety [90] days); AND
- The patient must be stable or experienced no more than one clinical relapse* in the past year; AND
- The patient has a recent EDSS score ≤ 5.

Dosage: 120 mg twice daily.
Maintenance: 240 mg twice daily

Duration of Approval: 1 year

*Renewal requests where patients have experienced more than one (1) clinical relapse in the past year are to be externally reviewed.

As applicable, please also include information regarding the requesting physician’s specialty (e.g. is the physician a neurologist or a physician with specialized experience with multiple sclerosis (MS), the name of the MS clinic where the patient was examined, or an MS consult note as this information may reduce the turnaround times for assessment.

Duration of Approval: First Renewal: 2 years
Second and subsequent renewals: 5 years
Fingolimod
Brand(s): Gilenya and generics
DOSAGE FORM/ STRENGTH: 0.5 mg capsule

As monotherapy for the treatment of patients with Relapsing Remitting Multiple Sclerosis (RRMS) who meet all of the following criteria:

- 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 attack(s), the date(s) of the attack(s), and the neurological findings; AND

- Failure to respond to full and adequate courses\(^1\) of at least one of interferon OR glatiramer acetate OR dimethyl fumarate; OR teriflunomide OR ocrelizumab OR documented intolerance or contraindication to 2 of the above listed therapies; AND

- Experienced one or more clinically disabling relapses in the previous year; AND

- Has had a 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.

- Is being followed by a neurologist experienced in the management of RRMS.

- Has a current EDSS of less than or equal to 5.5 (i.e. patients must be able to ambulate at least 100 meters without assistance).

Exclusion Criteria (Patients meeting any of the following exclusion criteria will not be funded):

- Patient’s receiving combination therapy of Gilenya with other disease modifying therapies (e.g. Aubagio, Avonex, Betaseron, Copaxone/Glatect, Extavia, Rebif, Extavia, Ocrevus, Tysabri, and Tecfidera).

- Patients with EDSS greater than 5.5

- Patients who have had a heart attack or stroke in the last 6 months of the funding request, history of sick sinus syndrome, atrioventricular block, significant QT prolongation, bradycardia, ischemic heart disease, or congestive heart failure.

- Patients younger than 18 years of age.

- Patients requesting Gilenya due to needle phobia or preference for oral therapy over injection who do not have a clinical contraindication to interferon or glatiramer therapy.

- Skin reactions at the site of injection do NOT qualify as a contraindication to interferon or glatiramer therapy.
Fingolimod
Brand(s): Gilenya
DOSAGE FORM/ STRENGTH: 0.5 mg capsule

Dosage: 0.5 mg once daily

Failure to respond to full and adequate courses: defined as having received a trial of at least 6 months of interferon or glatiramer or dimethyl fumarate therapy or teriflunomide AND experienced at least one disabling relapse (attack) while on interferon or glatiramer or dimethyl fumarate or teriflunomide.

MRI reports do NOT need to be submitted with the initial request.

Duration of Approval: 1 year

Renewals are considered. Renewals can be submitted through the Telephone Request Service and will be considered for patients who have benefited from therapy.

Physicians must provide the following information:

- Documentation providing the date and details of the Patient’s most recent neurological examination and EDSS scores (exam must have occurred within the last ninety (90) days); AND

- Evidence that the patient is stable and has experienced no more than one (1) disabling attack/relapse in the past year. (Note: If the Patient has had more than one attack/relapse, the request will be sent for external review. Please include details of the attack(s) including the dates on which they occurred); AND

- A recent Expanded Disability Status Scale (EDSS) that is less than or equal to 5.5 (Note: Requests with an EDSS greater than 5.5 will not be funded).

Dosage: 0.5 mg once daily.

Duration of Approval: First Renewal: 2 years

Second and subsequent renewals: 5 years
Glatiramer acetate – See Formulary listing for Glatect

Brand(s): Copaxone, Glatect (available as Limited Use drug on ODB formulary)

DOSAGE FORM/ STRENGTH: 20 mg/mL pre-filled syringe for subcutaneous injection

Effective September 27, 2018, Glatiramer as Copaxone for the treatment of Relapsed Refractory Multiple Sclerosis (RRMS)/ Clinically Definite Multiple Sclerosis (CDMS) or Clinically Isolated Syndrome (CIS) will only be considered for funding for existing EAP renewals. All EAP requests for patients treatment-naïve to Copaxone should consider Glatiramer as Glatect upon meeting Limited Use Criteria on the Ontario Drug Benefit Formulary effective on September 27, 2018.

For the treatment of Clinically Definite Multiple Sclerosis (CDMS) or Clinically Isolated Syndrome (CIS) (see criteria in next section. Also, note that patients who are treatment naïve to Copaxone should refer to the formulary for consideration of Glatect.

For CDMS: Copaxone requests for patients with CDMS will be reviewed by external medical experts when the following information is provided:

- Date and details of the most recent neurological examination (within the last 90 days); and
- Dates and details (e.g., neurological findings) of at least two clinical attacks, including one clinical attack within the past year; and
- EDSS score ≤ 5.

Duration of Approval: 1 year

Renewal requests for Copaxone can be submitted through the Telephone Request Service and will be considered for patients who have benefited from therapy and have an EDSS score ≤ 6.

The physician must provide the following information:

- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- The patient’s most recent EDSS score.

As applicable, include information regarding the requesting physician’s specialty (e.g. is the physician a neurologist or a physician with specialized experience with multiple sclerosis (MS), the name of the MS clinic where the neurologist is based, or an MS consult note supporting the diagnosis as this information may reduce the turnaround times for assessment.

Duration of Approval: First Renewal: 2 years
Second and subsequent renewals: 5 years

**Interferon beta-1a**

**Brand(s):** Avonex PS, Avonex Pen  
**DOSAGE FORM/ STRENGTH:** 30 mcg/ 0.5mL prefilled syringe for intramuscular injection, 30 mcg single-use prefilled auto-injector

For the treatment of Clinically Definite Multiple Sclerosis (CDMS) or Clinically Isolated Syndrome (CIS) (see CIS criteria in next section).

For CDMS: Avonex requests for patients with CDMS will be reviewed by external medical experts when the following information is provided:

- Details of the most recent neurological examination within the last ninety (90) days, including a description of any recent attacks (date and neurological findings)
- The patient has experienced at least two clinical attacks including one clinical attack within the past year
- MRI findings as applicable
- The patient’s EDSS is less than or equal to 6.0

**Renewal** requests for Avonex can be submitted through the Telephone Request Service. Avonex renewals will be considered for patients who have benefited from therapy. Patients must be stable (i.e. no relapses or attacks during the last year) and the patient’s EDSS must be less than or equal to 6.0

The physician must provide the following information:

- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- The patient’s most recent EDSS score.

As applicable, include information regarding the requesting physician’s specialty (e.g. is the physician a neurologist or a physician with specialized experience with multiple sclerosis (MS), the name of the MS clinic where the neurologist is based, or an MS consult note supporting the diagnosis as this information may reduce the turnaround times for assessment.

**Duration of Approval:** First Renewal: 2 years  
Second and subsequent renewals: 5 years
Interferon beta-1a
Brand(s): Rebif
DOSAGE FORM/ STRENGTH: 22 mcg and 44 mcg prefilled syringe for subcutaneous injection, 66 mcg/ml and 132 mcg/ml pre-filled cartridge

For the treatment of Clinically Definite Multiple Sclerosis (CDMS) or Clinically Isolated Syndrome (CIS) (see CIS criteria in next section).

For CDMS: Rebif requests for patients with CDMS will be reviewed by external medical experts when the following information is provided:

- Date and details of the most recent neurological examination (within the last 90 days); and
- Dates and details (e.g., neurological findings) of at least two clinical attacks, including one clinical attack within the past year; and
- EDSS score ≤ 6.

Duration of Approval: 1 year

Renewal requests for Rebif can be submitted through the Telephone Request Service and will be considered for patients who have benefited from therapy and have an EDSS score ≤ 6. The physician must provide the following information:

- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- The patient’s most recent EDSS score.

As applicable, include information regarding the requesting physician’s specialty (e.g. is the physician a neurologist or a physician with specialized experience with multiple sclerosis (MS), the name of the MS clinic where the neurologist is based, or an MS consult note supporting the diagnosis as this information may reduce the turnaround times for assessment.

Duration of Approval: First Renewal: 2 years

Second and subsequent renewals: 5 years
Interferon beta-1b
Brand(s): Betaseron
DOSAGE FORM/ STRENGTH: 0.3 mg/vial subcutaneous injection

For the treatment of Clinically Definite Multiple Sclerosis (CDMS) or Clinically Isolated Syndrome (CIS) (see criteria in next section)

For CDMS: Betaseron requests for patients will be reviewed by external medical experts when the following information is provided:

- Date and details of the most recent neurological examination (within the last 90 days); AND
- Dates and details (e.g., neurological findings) of at least two clinical attacks, including one clinical attack within the past year; AND
- EDSS score ≤ 6.

Duration of Approval: 1 year

Renewal requests for Betaseron can be submitted through the Telephone Request Service and will be considered for patients who have benefited from therapy and have an EDSS score ≤ 6.

The physician must provide the following information:

- Description of the patient’s clinical course in the last year, including details of all attacks; AND
- Date and details of the most recent neurological examination (within the last 90 days); AND
- The patient’s most recent EDSS score.

As applicable, include information regarding the requesting physician’s specialty (e.g. is the physician a neurologist or a physician with specialized experience with multiple sclerosis (MS), the name of the MS clinic where the neurologist is based, or an MS consult note supporting the diagnosis as this information may reduce the turnaround times for assessment.

Duration of Approval: First Renewal: 2 years

Second and subsequent renewals: 5 years
Interferon beta-1b
Brand(s): Extavia
DOSAGE FORM/ STRENGTH: 0.3 mg/vial subcutaneous injection

For the treatment of Clinically Definite Multiple Sclerosis (CDMS) or Clinically Isolated Syndrome (CIS) (see criteria in next section).

For CDMS: Extavia requests for patients will be reviewed by external medical experts when the following information is provided:

- Date and details of the most recent neurological examination (within the last 90 days) AND
- Dates and details (e.g., neurological findings) of at least two clinical attacks, including one clinical attack within the past year AND
- EDSS score ≤ 6.

Duration of Approval: 1 year

Renewal requests for Extavia can be submitted through the Telephone Request Service and will be considered for patients who have benefited from therapy and have an EDSS score ≤ 6.

The physician must provide the following information:

- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- The patient’s most recent EDSS score.

As applicable, include information regarding the requesting physician’s specialty (e.g. is the physician a neurologist or a physician with specialized experience with multiple sclerosis (MS), the name of the MS clinic where the neurologist is based, or an MS consult note supporting the diagnosis as this information may reduce the turnaround times for assessment.

Duration of Approval: 2 years

Second and subsequent renewals: 5 years
Modafinil
Brand(s): Alertec
DOSAGE FORM/ STRENGTH: 100 mg tablet

For the treatment of fatigue in patients with multiple sclerosis who have demonstrated a lack of response to or an inability to tolerate amantadine.

Note: See additional indications and criteria under “CNS” drugs

Duration of Approval: Lifetime

Natalizumab
Brand(s): Tysabri
DOSAGE FORM/ STRENGTH: 300 mg/15 mL concentrate for solution for intravenous infusion

Initiation Criteria:

As monotherapy for the treatment of Rapidly Evolving Severe Relapsing-Remitting Multiple Sclerosis (RES-RRMS) for the patient who meets all the following:

a) 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, including the corresponding dates, and the neurological findings; AND

b) Has been diagnosed with MS; AND

c) Is 18 to 65 years of age; AND

d) Has a current EDSS of less than or equal to 5.0; AND

e) 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 (i.e. 3 or more new lesions); 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;

f) has failed to respond to full and adequate courses of at least one of interferon OR glatiramer acetate OR dimethyl fumarate; OR teriflunomide OR documented intolerance or contraindication to 2 of the 3 therapies. (Note that needle phobia is not acceptable.)
Natalizumab
Brand(s): Tysabri
DOSAGE FORM/STRENGTH: 300 mg/15 mL concentrate for solution for intravenous infusion

g) is being followed by a neurologist experienced in the management of RRMS

h) details of past treatment, including dates and Patient response;

¹Failure to respond to a full and adequate course: defined as a trial of at least 6 months of interferon or glatiramer therapy or dimethyl fumarate AND experienced at least one disabling relapse (attack) while on interferon or glatiramer or dimethyl fumarate.

MRI reports do NOT need to be submitted with the initial request.

Duration of Approval: 1 year

Renewals will be considered for requests meeting the following;

(a) Documentation providing the date and details of the patient’s most recent neurological examination and EDSS scores (exam must have occurred within the last ninety (90) days); AND

(b) Evidence that the Patient is stable and has experienced no more than one (1) disabling attack/relapse in the past year (Note: if the Patient has had more than one attack/relapse, the request will be sent for external review); AND

(c) A recent Expanded Disability Status Scale (EDSS) that is less than or equal to 5.0 (Note that the request will be rejected if the EDSS is greater than 5.0).

Duration of Approval: First Renewal: 2 years

Second and subsequent renewals: 5 years
Initiation Criteria

For treatment of Early Primary Progressive Multiple Sclerosis (PPMS) in adult patients who meet ALL of the following criteria:

1. 18 years of age or older;
2. Diagnosis of early PPMS is confirmed based on McDonald 2017 diagnostic criteria meeting the following;
   i) Patient has had one year of disability progression (retrospectively or prospectively determined) independent of clinical relapse; AND
   ii) Two or more of the following;
       • One or more T2-hyperintense lesions (symptomatic or asymptomatic) characteristic of multiple sclerosis in one of more of the following brain regions periventricular, cortical, juxtacortical or infratentorial
       • Two or more T2-hyperintense lesions (symptomatic or asymptomatic) in the spinal cord
       • Presence of CSF-specific oligoclonal bands
3. Level of disability from disease meeting the below:
   i) A recent Expanded Disability Status Scale (EDSS)\(^1\) score between 3.0 and 6.5 prior to initiation of ocrelizumab; AND
   ii) A Functional Systems Scale (FSS) score of at least 2.0 for the pyramidal system due to lower extremity findings (Note that FSS scores associated with disability in other systems such as brainstem or cerebellar can be considered);
4. Disease duration from onset of multiple sclerosis meeting one of the below:
   i) Less than 15 years for those with an EDSS score greater than 5.0
   ii) Less than 10 years for those with an EDSS score equal to or less than 5.0
   iii) PPMS has been progressive in the last 3 years in the absence of activity
5. The drug request is from a neurologist experienced in the management of PPMS from one of the MS Society recognized Ontario MS clinics\(^2\) or includes a consult note from a neurologist from one of these clinics supporting the diagnosis and the treatment with ocrelizumab.

\(^1\)A “recent” score is an EDSS evaluated within the prior 6 months. Consideration will be provided for results from a neurological exam within the prior 12 months upon confirmation that the patient’s clinical status has not deteriorated.
Ocrelizumab
Brand(s): Ocrevus
DOSAGE FORM/ STRENGTH: 300 mg Injection

2MS Society recognized Ontario MS clinics*:

Hamilton MS Clinic HHS, McMaster University
Kingston MS Clinic, Kingston General Hospital
London MS Clinic, London Health Sciences Centre
Ottawa MS Research clinic, Ottawa Hospital General Campus
Ottawa Pediatric MS Clinic, CHEO
Toronto MS Clinic, St Michael’s Hospital
Toronto Pediatric MS Clinic, The Hospital for Sick Children
Sunnybrook Health Sciences Centre

*Note: Requests for patients who is under the care of a community neurologist working outside of one of the MS Society recognized Ontario MS clinics can be considered on a case-by-case basis.

Exclusion Criteria:

Patients with an EDSS score equal to or greater than 7.0

Dosage: Initial dose of 300 mg intravenous infusion, followed 2 weeks later by a second 300 mg intravenous infusion. Subsequent doses of single 600 mg intravenous infusion every 6 months after the first initial dose.

Renewal Criteria:

Ongoing funding will be provided for those who continue to benefit from treatment and who have an Expanded Disability Status Scale (EDSS) score less than 7.0.

Duration of Approval of Initials and Renewals: 18 months
Initiation Criteria:

**For the treatment of Relapsing Remitting Multiple Sclerosis (RRMS) in adult patients with active disease meeting ALL the following criteria:**

1. 18 years of age or older
2. Diagnosis of RRMS is in accordance with the McDonald 2017 criteria demonstrating dissemination of lesions in the central nervous system in space and time meeting the following;
   - 2 or more attacks¹ and clinical evidence of 2 or more lesions²; OR
   - 2 or more attacks and clinical evidence of 1 lesion with clear historical evidence of prior attack involving lesion in different location;

¹If the patient has experienced only one attack, the patient must meet ONE of the additional criteria of dissemination in time in the list below:
   - Additional clinical attack
   - Simultaneous presence of both enhancing and non-enhancing, symptomatic or asymptomatic MS-typical MRI lesions; OR new T2 or enhancing MRI lesion compared to baseline scan (without regard to timing of baseline scan)
   - Presence of cerebrospinal fluid (CSF)-specific oligoclonal bands

²If the patient has evidence of only one lesion the patient must meet ONE of the additional criteria of dissemination in space in the list below:
   - additional clinical attack implicating different CNS site
   - 1 or more MS-typical T2 lesions in 2 or more areas of the Central Nervous System (CNS): periventricular, cortical, juxtacortical, infratentorial or spinal cord
   AND
3. Patient has experienced a clinical relapse and/or new MS lesions in the last 2 years; AND
4. Patient has an EDSS score less than 6.0 before start of therapy; AND
5. Ocrelizumab is used as monotherapy; AND
6. The drug request is from a neurologist experienced in the management of RRMS from one of the MS Society recognized Ontario MS clinics³ or includes a consult note from a neurologist from one of these clinics supporting the diagnosis.
Ocrelizumab
Brand(s): Ocrevus
DOSAGE FORM/ STRENGTH: 300 mg Injection

3MS Society recognized Ontario MS clinics*:

Hamilton MS Clinic HHS, McMaster University
Kingston MS Clinic, Kingston General Hospital
London MS Clinic, London Health Sciences Centre
Ottawa MS Research clinic, Ottawa Hospital General Campus
Ottawa Pediatric MS Clinic, CHEO
Toronto MS Clinic, St Michael’s Hospital
Toronto Pediatric MS Clinic, The Hospital for Sick Children
Sunnybrook Health Sciences Centre

*Note: Requests for patients who is under the care of a community neurologist working outside of one of the MS Society recognized Ontario MS clinics can be considered on a case-by-case basis.

Exclusion criteria:
1. Combination therapy with another disease modifying therapy for RRMS will not be reimbursed.
2. Patients with an EDSS score equal to or greater than 7.0

Dosage: Initial dose of 300 mg intravenous infusion, followed 2 weeks later by a second 300 mg intravenous infusion. Subsequent doses of single 600 mg intravenous infusion every 6 months after the first dose.

Renewal Criteria:

Ongoing funding will be provided for those who continue to benefit from treatment and who have an Expanded Disability Status Scale (EDSS) score less than 7.0

When requesting renewal of funding, information that should be provided should include:

- Date and details of the most recent neurological examination and EDSS scores to support ongoing benefit from therapy.
- Clinical details of the date and onset of clinical attacks/relapses
- Information to support that the patient is stable/not demonstrating a sub-optimal response

Renewal requests where patients have experienced more than 1 attack in the past year will be externally reviewed.

Approval Duration of Initial and Renewals: 18 months
For the treatment of Clinically Definite Multiple Sclerosis (CDMS)/ Relapsing remitting multiple sclerosis (RRMS) in patients meeting the following criteria:

Plegridy requests will be reviewed by external medical experts when the following information is provided:

- Details of the most recent neurological examination within the last ninety (90) days, including a description of any recent attacks (date and neurological findings)
- The patient has experienced at least two clinical attacks in his or her lifetime, including one clinical attack within the past 12 months preceding the EAP request;
- MRI findings as applicable
- The patient’s EDSS is less than or equal to 6.0

Duration of Approval: 1 year

Renewal requests for Plegridy can be submitted through the Telephone Request Service. Plegridy renewals will be considered for patients who have benefited from therapy. Patients must be stable (i.e. no relapses or attacks during the last year) and the patient’s EDSS must be less than or equal to 6.0.

The physician must provide the following information:

- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- The patient’s most recent EDSS score.

*Renewal requests where patients have experienced more than one (1) clinical relapse in the past year will be considered on a case-by-case basis through an external review.

As applicable, include information regarding the requesting physician’s specialty (e.g. is the physician a neurologist or a physician with specialized experience with multiple sclerosis (MS), the name of the MS clinic where the neurologist is based, or an MS consult note supporting the diagnosis as this information may reduce the turnaround times for assessment.

Duration of Approval: First Renewal: 2 years
Second and subsequent renewals: 5 years

**Teriflunomide**

*Brand(s): Aubagio*

**DOSAGE FORM/ STRENGTH:** 14 mg tablet

**For the treatment of relapsing-remitting multiple sclerosis (RRMS)** in patients who meet the following criteria;

i) the physician making the request on behalf of the patient is a neurologist who is experienced in the management of RRMS; AND

ii) the physician provides documentation of the patient’s most recent neurological examination which must have been conducted within ninety (90) days preceding the submission of the EAP request. This must include a description and dates of any recent attacks and other pertinent neurological findings; AND

iii) the patient’s diagnosis is confirmed to be RRMS; AND

xi) the patient has experienced one or more clinical attacks/relapses in the year preceding the request; AND

xii) the patient has a recent Expanded Disability Status Scale (EDSS) score that is equal to or less than 5.0 prior to starting therapy with teriflunomide.

Dosage: 14 mg once daily

**Duration of Approval:** 1 year

**Renewals** for the funding of teriflunomide will be considered in patients who meet the following criteria;

i) the physician provides documentation of the date and details of the patient’s most recent neurological examination and EDSS scores (the examination must have occurred within the last ninety [90] days preceding the submission of the renewal request); AND

ii) the physician confirms that the Patient is stable and has experienced no more than one (1) clinical relapse in the past year¹; AND

iii) the patient’s most recent EDSS score while on teriflunomide is less than or equal to 5.0.

¹Renewal requests where the patient has experienced more than 1 clinical relapse in the past year will be considered on a case-by-case basis with the assistance of external medical consultants.

Dosage: 14 mg once daily.
Duration of Approval 2 years
# CLINICALLY ISOLATED SYNDROME DRUGS

<table>
<thead>
<tr>
<th>Drug</th>
<th>Brand(s)</th>
<th>DOSAGE FORM/ STRENGTH</th>
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<tbody>
<tr>
<td>Glatiramer acetate</td>
<td>Copaxone</td>
<td>20 mg/mL pre-filled syringe for subcutaneous injection</td>
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<tr>
<td>Interferon beta-1a</td>
<td>Avonex PS, Avonex Pen</td>
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<td>Betaseron</td>
<td>0.3 mg/vial subcutaneous injection</td>
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<tr>
<td></td>
<td>Extavia</td>
<td>0.3 mg/vial subcutaneous injection</td>
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For the treatment of Clinically Isolated Syndrome (CIS): requests for patients who have experienced a single demyelinating event will be reviewed by external medical experts when the following information is provided:

- Date and details of the most recent neurological examination which must have been conducted within the last ninety days of the request;
- The patient’s EDSS is less than or equal to 6.0 (please provide EDSS score); AND
- The patient’s clinically isolated syndrome occurred within the last twelve months.

**Duration of Approval:** 1 year

**Renewal** requests will be assessed according to the following criteria:

- the requesting physician provides the date and details of the patient’s most recent neurological examination and EDSS scores;
- the patient’s neurological examination occurred within that last ninety days;
- the patient is stable (i.e. no relapses or attacks during the last year) and
- the patient’s EDSS is less than or equal to 6.0
OCULAR TREATMENTS

Tocilizumab
Brand(s): Actemra
DOSAGE FORM/ STRENGTH: 162 mg/0.9 mL inj (PFS), 162mg/0.9mL Auto Inj.

For the treatment of new onset or relapsed Giant Cell Arteritis (GCA) in adult patients meeting all the following criteria,

- Symptomatic for GCA; AND
- Diagnosis of GCA confirmed by temporal artery biopsy and/or imaging tests (i.e. magnetic resonance angiography, computed tomography angiography or positron emission scanning)\(^1\); AND
- Tocilizumab subcutaneous is used as combination therapy with 20 mg to 60 mg of prednisone (or an equivalent corticosteroid) with subsequent corticosteroid tapering as symptoms stabilize; AND
- Prescribed by a rheumatologist or a prescriber with expertise in the diagnosis and management of GCA.

\(^1\)Where these tests are not available or where a result may be deemed unreliable (e.g. a negative biopsy in a patient on corticosteroids), the prescriber may C-reactive protein and/or Erythrocyte Sedimentation Rate results with the request.

Recommended dose:

162 mg sc once a week (or once every other week, based on clinical considerations) in combination with a tapering course of corticosteroid.

Approval Duration: 1 year

Renewals will be considered on a case-by-case basis.

Approval Duration of renewals: 1 year
Infliximab - See formulary for funded biosimilars
Brand(s): Remicade and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 100 mg/Vial Injection for infusion

For the treatment of severe non-infectious ocular inflammatory disease (OID) in patients meeting one of the following criteria;

- Experienced failure, intolerance, or contraindication to oral corticosteroid (or topical corticosteroid for anterior uveitis) and failure or intolerance to at least one immunosuppressive therapy; OR

- For the treatment of chronic Juvenile Idiopathic Arthritis (JIA)-associated uveitis after failure or intolerance to a first-line immunosuppressive agent; OR

- For patients who have immediately vision-threatening OID and do not meet the above criteria, where consultation notes/letter from an ophthalmologist expert specializing in OIDs (who may be the requesting physician) confirm the severity of the patient’s condition and indicate detailed rationale for an immediate biologic therapy (e.g. ocular inflammation associated with Behcet’s disease; severe non-necrotizing scleritis; necrotizing scleritis; etc.); AND

- Patient must be followed by a uveitis specialist, a retina specialist familiar with ocular inflammatory diseases, or a pediatric ophthalmologist.

**Approved Dose:** Infliximab 5-10 mg/kg IV at weeks 0, 2, 6 and maintenance every 4-8 weeks

**Duration of Approval:** 1 year

**Renewals** will be considered for requests where consultation notes or a letter is provided by the requesting physician to confirm that treatment has resulted in improvement/stability of vision and other treatment goals (e.g., remission from/control of ocular inflammation) have been met.

**Duration of Approval:** 2 years
Adalimumab – See Formulary for funded biosimilars
Brand(s): Humira and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 40 mg/0.8 mL prefilled syringe, 40mg/0.8mL and 20 mg/0.2 mL prefilled pens for subcutaneous injection

- Experienced failure, intolerance, or contraindication to oral corticosteroid (or topical corticosteroid for anterior uveitis) and failure or intolerance to at least one immunosuppressive therapy; OR

- For the treatment of chronic Juvenile Idiopathic Arthritis (JIA)-associated uveitis after failure or intolerance to a first-line immunosuppressive agent; OR

- For patients who have immediately vision-threatening OID and do not meet the above criteria, where consultation notes/ letter from an ophthalmologist expert specializing in OIDs (who may be the requesting physician) confirm the severity of the patient’s condition and indicate detailed rationale for an immediate biologic therapy (e.g. ocular inflammation associated with Behcet’s disease; severe non-necrotizing scleritis; necrotizing scleritis; etc.); AND

- Patient must be followed by a uveitis specialist, a retina specialist familiar with ocular inflammatory diseases, or a pediatric ophthalmologist.

Approved Dose: Adalimumab 40 mg subcutaneous every 1 to 2 weeks.

Duration of Approval: 1 year

Renewals will be considered for requests where consultation notes or a letter is provided by the requesting physician to confirm that treatment has resulted in improvement/stability of vision and other treatment goals (e.g., remission from/control of ocular inflammation) have been met.

Duration of Approval: 2 years
Rituximab
Brand(s): Riximyo, Ruxience, and Truxima (biosimilar); Rituxan (biologic originator)
DOSAGE FORM/ STRENGTH: 10 mg/mL intravenous injection

For the treatment of severe non-infectious ocular inflammatory disease (OID) in patients failed or did not tolerate treatment with infliximab or adalimumab; OR has contraindication to anti-TNF therapy AND who meet one of the following criteria:

- Experienced failure, intolerance, or contraindication to oral corticosteroid (or topical corticosteroid for anterior uveitis) and failure or intolerance to at least one immunosuppressive therapy; OR

- For the treatment of chronic Juvenile Idiopathic Arthritis (JIA)-associated uveitis after failure or intolerance to a first-line immunosuppressive agent; OR

- For patients who have immediately vision-threatening OID and do not meet the above criteria, where consultation notes/ letter from an ophthalmologist expert specializing in OIDs (who may be the requesting physician) confirm the severity of the patient’s condition and indicate detailed rationale for an immediate biologic therapy (e.g. ocular inflammation associated with Behcet’s disease; severe non-necrotizing scleritis; necrotizing scleritis; etc.); AND

- Patient must be followed by a uveitis specialist, a retina specialist familiar with ocular inflammatory diseases, or a pediatric ophthalmologist.

**Approved Dose:** Rituximab up to 1000 mg IV per infusion at days 1 & 15 and 3rd infusion at 6-12 months.

Note that maintenance rituximab infusions are not funded.

**Duration of Approval:** 1 year

**Renewals** will be considered for requests where;

- Consultation notes or a letter is provided by the requesting physician to confirm that treatment has resulted in improvement/stability of vision and other treatment goals (e.g., remission from/control of ocular inflammation) have been met; AND

Patients must also have demonstrated subsequent deterioration of symptoms, at least 6 months from the last dose of rituximab.

**Duration of Approval:** 2 years
Cyclosporine 0.1% eye drops
Brand(s): Verkazia
DOSAGE FORM/ STRENGTH: 0.1% eye drops
Effective date: March 3, 2021

Initiation criteria:

For the treatment of severe vernal keratoconjunctivitis (VKC) in patients who meet ALL the following criteria:

- Patient is between 4 and 18 years of age inclusive; AND
- Diagnosis of severe VKC as defined by one of the following evaluations of severity:
  - Grade 3 (severe) or 4 (very severe) on the Bonini Scale; OR
  - Grade 4 (marked) or 5 (severe) on the modified Oxford Scale; AND
- Documentation detailing the patient’s baseline severity of signs and symptoms of VKC is provided prior to treatment initiation with Verkazia; AND
- Patient is under the care of a physician with expertise in the diagnosis and management of VKC.

Note: Patients previously treated with cyclosporine 0.1% but who discontinued treatment upon resolution of VKC signs and symptoms are eligible to reinitiate treatment if signs and symptoms of severe VKC recur and they meet the first and second initiation criteria.

Initial approval period: 6 months

Exclusion criteria:

Verkazia will not be funded in patients meeting ANY of the following criteria:

- Patient is older than 18 years of age;
- Patient has asymptomatic, pre-symptomatic, or moderate VKC;
- Patients with active or suspected ocular or peri-ocular infection;
- Patients with ocular malignancies or premalignant conditions.

Discontinuation Criteria:

Verkazia will not be funded upon meeting ONE or more of the following;

- There is no demonstrated observable improvement in the patient’s VKC signs and symptoms after 4 months of Verkazia therapy (compared to baseline); OR
- The patient’s VKC signs and symptoms have been resolved.

Case-by-case consideration may be provided for patients who have responded to therapy but who continue to experience recurrent VKC symptoms
Abiraterone
Brand(s): Zytiga and generics (see formulary for full list of funded generics)
DOSAGE FORM/ STRENGTH: 250 mg, 500 mg tablet

Reimbursement criteria for Zytiga in patients who have not trialed docetaxel.

For the treatment of metastatic castrate-resistant prostate cancer (mCRPC) in patients who meet the following criteria:

- Zytiga is being used in combination with prednisone; AND
- The patient is asymptomatic or mildly symptomatic after failure of androgen deprivation therapy; AND
- The patient has an ECOG* ≤ 1; AND
- The Patient must not meet any of the exclusion\(^1\) criteria stated below.

*ECOG = Eastern Cooperative Oncology Group Status

(Please provide clinical information as objective evidence that the above criteria are met (e.g. castrate testosterone level, prostate surface antigen levels, evidence of metastatic disease such as presence and location of lesions, surgical procedures related to the condition, and name(s), date, duration of androgen deprivation therapy used details of the response to therapy, labwork or clinical confirmation to support that the patient does not meet any of the exclusion criteria.)

Approved dosage: 1000 mg once daily will be funded until there is evidence of disease progression.

Duration of Approval: 1 year

Renewals will be considered in patients with evidence of not having had disease progression while on Zytiga therapy.

\(^1\)Exclusion Criteria:

Funding for Zytiga will NOT be approved in patients who meet any ONE (or more) of the following exclusion criteria:

- the Patient has viral hepatitis or chronic liver disease; OR
- the Patient has clinically significant heart disease; OR
- Zytiga is being prescribed for combination use with Jevtana or Xtandi for mCRPC; OR
- The patient has received prior chemotherapy for mCRPC.
Abiraterone
Brand(s): Zytiga and generics (see formulary for full list of funded generics)
DOSAGE FORM/ STRENGTH: 250 mg, 500 mg tablet

Reimbursement criteria for Zytiga in patients who are requesting Zytiga after a trial of docetaxel.

For the treatment of metastatic castrate-resistant prostate cancer (mCRPC) in patients who meet the following criteria:

- Zytiga is being used in combination with prednisone; AND
- The patient’s cancer has progressed after having received prior docetaxel containing therapy; AND
- The patient has ECOG* ≤ 2.
- Patients must not meet ANY of the exclusion criteria for funding stated below.

*ECOG = Eastern Cooperative Oncology Group Status

Requests for patients who initiated Jevtana (cabazitaxel) or Xtandi (enzalutamide) therapy within the three (3) months preceding the EAP request for Zytiga and who have not had disease progression, will be considered on a case-by-case basis.

Approved dosage: 1000 mg once daily will be funded until there is evidence of disease progression.

Renewals will be considered in patients with evidence of not having had disease progression while on Zytiga therapy.

Exclusion Criteria:

Funding for Zytiga will NOT be approved in patients who meet any ONE (or more) of the following exclusion criteria:

- the Patient has viral hepatitis or chronic liver disease; OR
- the Patient has clinically significant heart disease; OR
- Zytiga is being prescribed for combination use with Jevtana or Xtandi for mCRPC; OR
- the Patient has already used Zytiga in the pre-docetaxel setting.

Duration of Approval: 1 year
For the treatment of adult patients with chronic lymphocytic leukemia (CLL) who meet the following criteria:

1. Diagnosed with active disease according to one or more of the International Workshop on Chronic Lymphocytic Leukemia (iwCLL) 2008 criteria; AND

2. Meets one of the following circumstances of use:
   i. First line use in a previously untreated patient who presents with one or more of the following cytogenic markers:
      • chromosome 17p deletion; OR
      • TP53 mutation, OR
      • unmutated immunoglobulin heavy chain variable region (IgHV); OR
   ii. Patient with relapsed or refractory CLL who has experienced disease progression on at least one prior systemic therapy/regimen.*

   AND

3. Acalabrutinib will be used as monotherapy; AND

4. Patient has good performance status.

*A prior line of therapy should include a fludarabine-based regimen in fit patients.

Renewal Criteria:

Renewals will be considered in patients until disease progression (as defined based on published iwCLL [2018]) or unacceptable toxicity.

Exclusion Criteria: (Patients meeting the below will not be funded.)

1. Patients who have experienced disease progression on another Bruton’s tyrosine kinase (BTK) inhibitor (e.g. ibrutinib) for the treatment of CLL.

2. Patients with prolymphocytic leukemia.

3. Patients with current or history of Richter's syndrome.

4. Patients with central nervous system (CNS) lymphoma or leukemia.
Acalabrutinib
Brand(s): Calquence
DOSAGE FORM/ STRENGTH: 100 mg Capsule

Notes:

1. The Ministry will fund only one line of treatment with a BTK inhibitor for CLL.

2. The Ministry will not fund idelalisib following progression on a BTK inhibitor.

3. Patients may switch to acalabrutinib from another BTK inhibitor (e.g. Ibrutinib) as long as they have not experienced disease progression on another BTK inhibitor. Please include the reasons for requesting the switch with the application (e.g. intolerances, contraindications, etc.).

4. Acalabrutinib may be considered in patients diagnosed with small lymphocytic lymphoma (SLL) on a case-by-case basis upon meeting the above treatment eligibility criteria for CLL.

Approved Dosage for Initials and Renewals: 100 mg orally twice daily

Duration for Approval of Initials and Renewals: 1 year
Afatinib
Brand(s): Giotrif
DOSAGE FORM/ STRENGTH: 20 mg, 30 mg, 40 mg tablet

Initial requests:

For the treatment of patients with advanced or metastatic non-small cell lung cancer (NSCLC) who meet the following criteria;

- Afatanib is being used as first line therapy; AND
- Afatanib is being used as monotherapy; AND
- Patient’s cancer is EGFR positive

Dose: 40 mg orally once daily

Exclusion Criteria:
- Patients with EGFR wild-type, negative, or unknown mutation.
- Afatinib will not be considered for funding in patients who have progressed on a prior EGFR TKI targeted therapy.
- Not funded for 2\textsuperscript{nd} or 3\textsuperscript{rd} line or maintenance NSCLC.

Notes:
- Patients should be assessed for disease status at least every two months. Afatinib may be continued until evidence of disease progression or development of unacceptable toxicity requiring discontinuation of afatinib.
- Patients who receive afatinib 1\textsuperscript{st} line are NOT eligible for erlotinib in the 2\textsuperscript{nd} or 3\textsuperscript{rd} line or maintenance NSCLC setting.
- Requests for afatinib for patients who have initiated another EGFR TKI therapy (i.e. Iressa [gefitinib]) in the first line setting and who have not had disease progression will be considered on a case-by-case basis.

Renewal requests will be considered based on the following;

Afatinib 40 mg once daily may be continued until evidence of disease progression or development of unacceptable toxicity at which point the drug should be discontinued. Patients should have their disease status assessed at least every two months.

Exclusion Criteria:
- Patients with EGFR wild-type, negative, or unknown mutation.
- Afatinib will not be considered for funding in patients who have progressed on a prior EGFR TKI targeted therapy. Not funded for 2\textsuperscript{nd} or 3\textsuperscript{rd} line or maintenance NSCLC.
Initial Criteria:

For the treatment of anaplastic lymphoma kinase (“ALK”) – positive locally advanced (not amenable to curative therapy) or metastatic non-small cell lung cancer (NSCLC) in patients meeting ALL the following criteria;

- Alectinib is used as first line treatment OR after experiencing disease progression or intolerance on crizotinib\(^1, 2, 3\); AND
- Alectinib is used as monotherapy; AND
- Patient has good performance status (ECOG ≤ 2).

\(^1\)Patients who have progressed during or following first-line therapy with alectinib are not eligible to receive alectinib as a subsequent-line therapy.

\(^2\)Time-limited funding will be considered case-by-case in patients with ALK-positive NSCLC who have progressed on chemotherapy and crizotinib OR crizotinib and an immune checkpoint inhibitor commenced prior to the public funding of alectinib.

\(^3\)Include details of the intolerance including the grade of toxicity and reasons why crizotinib was not able to be used.

Exclusion criteria:

- Alectinib will not be funded if the patient has experienced disease progression while on an ALK inhibitor other than crizotinib.
- Alectinib will not be funded beyond third line.

Public funding will be considered for only one of Alectinib (Alecensaro) OR Ceritinib (Zykadia) and vice versa.

Recommended dose: 600 mg twice daily

Renewal Criteria:

Ongoing funding will be considered in patients who have not experienced disease progression or unacceptable toxicities to treatment with Alectinib.

Approval duration of initial and renewal requests: 1 year
Initiation Criteria for metastatic castration sensitive prostate cancer (mCSPC):

For the treatment metastatic castration sensitive prostate cancer (mCSPC) in patients who meet all of the following criteria:

1. Apalutamide is used in combination with androgen deprivation therapy (ADT)\(^1\); AND
2. Metastatic lesions detected on technetium-99m bone scan, computed tomography (CT), and/or magnetic resonance imaging (MRI); AND
3. Castration sensitive as defined by the patient being treatment naïve to an ADT OR ADT initiated within the prior 6 months before start of therapy with apalutamide OR ADT used in the neoadjuvant or adjuvant setting where the patient has been off treatment for 12 months or more prior to start of apalutamide; AND
4. Has not experienced disease progression with another androgen receptor axis targeted therapy (ARAT) for castration sensitive prostate cancer; AND
5. Patient has good performance status.

\(^1\)ADT is not required for patients with bilateral orchiectomy

Definitions for the purpose of the EAP funding criteria:

**ADT** – A first generation androgen deprivation therapy (e.g. goserelin, leuprolide, triptorelin, buserelin, degarelix)

**ARI** - A second generation androgen receptor inhibitor (e.g. apalutamide, darolutamide, enzalutamide)

**ARAT**- An androgen receptor axis targeted therapy (e.g. abiraterone, apalutamide, darolutamide, enzalutamide)

The following baseline levels are to be provided with the initial application:

i) Number of metastatic lesions on bone scan and in soft tissues

ii) Testosterone level

iii) Baseline (pre-treatment) PSA level

iv) Pre-treatment Gleason score (optional)
Apalutamide
Brand(s): Erleada
DOSAGE FORM/ STRENGTH: 60 mg Capsule

Notes:

1. Patients who have previously progressed on an ARI for prostate cancer will not be eligible for apalutamide in mCSPC. The Ministry will fund only one of apalutamide or enzalutamide in patients with mCSPC.
2. Patients treated with enzalutamide or darolutamide as part of a clinical trial may be eligible for apalutamide and will be considered on a case-by-case basis.
3. Patients who are currently on a treatment regimen with an ARAT for prostate cancer, must meet the initiation criteria if they wish to switch to publicly funded apalutamide for mCSPC.
4. Time limited funding consideration will be provided on a case-by-case basis for those patients who are using docetaxel in combination with ADT as long as there has been no progression and the treatment regimen has not been used for more than 6 months.

Approved Dosage for Initials and Renewals: 240 mg administered orally once daily.

Renewal Criteria:

Renewals will be considered in patients until disease progression (i.e. PSA elevation in addition to radiographic disease progression, or PSA progression in addition to clinical symptoms associated with cancer progression), development of castration resistant disease, or experiencing unacceptable toxicity while on apalutamide.

Exclusion Criteria: (Patients meeting any of the following will not be funded.)

1. Patients who have previously experienced disease progression on apalutamide or another ARI used in the setting of prostate cancer.
2. Apalutamide will not be funded as combination therapy with another ARAT.

Duration of initial and renewal approvals: 1 year
Apalutamide
Brand(s): Erleada
DOSAGE FORM/ STRENGTH: 60 mg Capsule

Initiation criteria for for Non-metastatic Castrate Resistant Prostate Cancer (nmCRPC):

For the treatment of high risk non-metastatic castration resistant prostate cancer (nmCRPC) in patients who meet all the following criteria:

1. Patient using apalutamide in combination with androgen deprivation therapy (ADT); AND
2. Has no detectable distant metastases as determined by CT, MRI, or technetium-99m bone scan; AND
3. Patient has castration resistant disease based on meeting all the following indicia observed while on continuous ADT treatment or post orchiectomy:
   a. Castrate serum testosterone levels: AND
   b. Biochemical progression defined as Three (3) prostate-specific antigen (PSA) rises at least 1 week apart, with the last PSA greater than 2ng/mL; and
4. Patient is at high risk for developing metastatic disease based on a Prostate-specific antigen doubling time (PSADT) of less than or equal to 10 months during continuous ADT; AND
5. Has Eastern Cooperative Oncology Group (ECOG) Performance Status less than or equal to 2.

Exclusion criteria: The following will not be reimbursed.

1. The patient received prior chemotherapy for the treatment of prostate cancer unless it was in the adjuvant or neoadjuvant setting.
2. The patient has previously experienced disease progression on prior treatment with enzalutamide or darolutamide used for prostate cancer.

Definitions for the purpose of the EAP funding criteria:

**ADT** – A first generation androgen deprivation therapy (e.g. goserelin, leuprolide, triptorelin, buserelin, degarelix).

**ARI** - A second generation androgen receptor inhibitor (e.g. apalutamide, darolutamide, enzalutamide).
Apalutamide
Brand(s): Erleada
DOSAGE FORM/ STRENGTH: 60 mg Capsule

Notes:
1. The Ministry will fund only one second generation androgen receptor inhibitor (ARI) in patients with non-metastatic castrate resistant prostate cancer.
2. Requests for apalutamide in patients who have initiated another ARI therapy in the nmCRPC setting and who have not experienced disease progression will be considered on a case-by-case basis.
3. Patients treated with an ARI as part of a clinical trial may be eligible for apalutamide and will be considered on a case-by-case basis.

Approved Dosage: 240 mg administered orally once daily.

Renewal Criteria:
Renewals will be considered in patients without evidence of radiographic disease progression or unacceptable toxicity while on apalutamide therapy.

Duration of initial and renewal approvals: 1 year
For the treatment of advanced renal cell carcinoma (MRCC) in patients who meet the following criteria:

1. Axitinib is being requested as treatment for a patient with advanced renal cell carcinoma in one of the following settings:
   
   i) Axitinib is being used in treatment naïve patients as first line therapy in combination with pembrolizumab in patients with disease in any risk category (Note that in this first line setting, cabozantinib may be funded as second line following progression to axitinib); OR
   
   ii) As second line therapy after failure to sunitinib or pazopanib in patients with disease in any risk category (Note that in this second line setting, only one of axitinib OR nivolumab OR cabozantinib will be funded); OR
   
   iii) As third line therapy in patients in the intermediate or poor risk disease category who have progressed on combination therapy with ipilimumab and nivolumab in first line and sunitinib or pazopanib used in second line. (Note that in this third line setting, only one of axitinib or cabozantinib will be funded.)

2. Patients must have good performance status.

NOTES:

- When used in combination with pembrolizumab in the first line setting, patients who are intolerant to pembrolizumab may stop pembrolizumab and continue treatment with axitinib as monotherapy until disease progression or development of unacceptable toxicities.
- When used in combination with pembrolizumab in the first line setting, patients experiencing disease progression or unacceptable toxicities may be considered for cabozantinib in second line, however, only two lines of treatment will be funded in this setting.
- Case-by-case consideration may be provided for patient with MRCC who have used and progressed on publicly funded older treatment regimens for MRCC such as everolimus, sorafenib, temsirolimus and/or interferons.

Funded Dosing Regimen: 5 mg twice a day. Dose may be adjusted based on individual response and tolerability.

Renewal Criteria: Renewals will be considered in patients who have not experienced disease progression or unacceptable toxicity while being treated with axitinib.

Approval duration of initials and renewals: 1 year
For the treatment of patients with Philadelphia chromosome-positive (Ph+) chronic myelogenous leukemia (CML) in patients meeting the following criteria;

1. 18 years of age or older; AND

2. Diagnosis of (Ph+) chronic phase, accelerated phase, or blast phase CML; AND

3. Bosutinib is used in one of the following clinical situations:

   i) As second line therapy after experiencing disease progression on imatinib, dasatinib, or nilotinib\(^1\) in first line: OR

   ii) as third line therapy after experiencing disease progression to imatinib, dasatinib, nilotinib\(^1\), or ponatinib in second line; OR

   iii) In patients who have not progressed on one or more prior TKIs but who have documented mutational drug resistance to imatinib, dasatinib, and/or nilotinib\(^1\) which make them clinically inappropriate treatment choices; OR

   iv) In patients who have not progressed on one or more TKIs but have experienced unacceptable intolerance or toxicity to one prior TKI (i.e. imatinib, dasatinib, or nilotinib\(^1\))

**Exclusion Criteria:**

1. Bosutinib will not be funded in combination with another oral TKI (e.g. imatinib, nilotinib, dasatinib, or ponatinib)

2. Bosutinib will not be funded as 4\(^{th}\) line treatment for CML or beyond.

\(^1\)Note that nilotinib is not funded in blast phase CML, therefore, considerations will only be applied for imatinib and dasatinib in patients with blast phase CML

**Renewal criteria:**

Renewal of funding will be considered upon confirmation from the patient’s clinician that the patient has experienced hematologic and/or cytogenic response and is expected to continue to do so

**Recommended Dosing:** 500mg per day

**Approval period for initials & renewals:** 1 year
Initiation Criteria for Advanced Renal Cell Carcinoma:

Cabozantinib (Cabometyx) will be reimbursed as monotherapy treatment of patients with advanced renal cell carcinoma (RCC) meeting one of the following situations:

1. As monotherapy, second line therapy in a Patient with any risk category (i.e. good, intermediate or poor risk) of advanced RCC after progression on at least one prior vascular endothelial growth factor receptor (VEGFR) tyrosine kinase inhibitor (TKI) therapy (e.g. Sunitinib, Pazopanib).

   If used in this second line setting, only one of cabozantinib or axitinib or nivolumab will be funded.

2. As monotherapy, third line therapy in a Patient with any risk category (i.e. good, intermediate or poor risk) of advanced RCC after progression on at least one prior vascular endothelial growth factor receptor (VEGFR) tyrosine kinase inhibitor (TKI) therapy (e.g. Sunitinib, Pazopanib) in first line and nivolumab in second line.

3. As monotherapy, third line therapy in a Patient with intermediate or poor risk advanced RCC after progression on an ipilimumab-nivolumab combination in first line and a vascular endothelial growth factor receptor (VEGFR) tyrosine kinase inhibitor (TKI) therapy (e.g. Sunitinib, Pazopanib) in second line.

   If used in this third line setting, only one of cabozantinib or axitinib will be funded.

Exclusion Criteria:

1. Patients who have experienced progression to cabozantinib for advance RCC will not be considered for EAP reimbursement for retreatment with cabozantinib in a subsequent line.

2. Cabozantinib will not be funded in combination with another treatment for advance RCC.

3. Cabozantinib will not be funded as first line therapy.

4. Cabozantinib will not be funded for patients when used as in fourth line or later therapy.
**Cabozantinib**

**Brand(s):** Cabometyx  
**DOSAGE FORM/ STRENGTH:** 20 mg, 40 mg, 60 mg tablet

Case-by-case consideration may be provided for patients who have experienced disease progression or intolerance to everolimus or temsirolimus or sorafenib used for advanced or metastatic RCC.

**Renewals** will be considered until clinically meaningful disease progression or the patient has experienced unacceptable toxicity.

Recommended dose: 60 mg daily.

Requests for 20mg and 40mg tablets should include reasons why the lower dosed tablets are required.

**Approval duration for initials and renewals:** 1 year

**Initiation Criteria for Advanced Hepatocellular Carcinoma:**

For the treatment of unresectable, advanced hepatocellular carcinoma (HCC) in adult patients meeting all of the following criteria prior to starting treatment with cabozantinib:

1. Patient is 18 years of age and older; **AND**
2. Cabozantinib will be used as monotherapy for HCC; **AND**
3. Cabozantinib will be used as second line therapy in a patient who has experienced disease progression during treatment with sorafenib OR lenvatinib for HCC; **AND**
4. Patient has good performance status with Eastern Cooperative Oncology Group (ECOG) Performance status less than or equal to 1; **AND**
5. Has a Child-Pugh class A liver function.

**Exclusion criteria:** Patients meeting any of the following criteria will not be funded:

1. Cabozantinib will not be funded in combination therapy with another therapy for HCC.
2. Cabozantinib will not be funded in patients with Child-Pugh class status of B or C.
3. Cabozantinib will not be funded as first-line therapy.
**Notes**

1. Only one of regorafenib or cabozantinib for the treatment of unresectable HCC will be funded in the second line setting.

2. Cabozantinib can be funded in patients who develop intolerances or toxicities to regorafenib and wish to switch to cabozantinib as long as they have not experienced disease progression on regorafenib and meet the initial funding criteria for cabozantinib.

Recommended dose: 60 mg orally once daily

Requests for 20mg and 40mg tablets should include reasons why the lower dosed tablets are required.

**Renewal criteria:**

Renewals will be considered until clinically meaningful disease progression\(^1\) or the patient has experienced unacceptable toxicity.

Please provide radiographic and/or scan results indicating no progression.

\(^1\) Evaluation according to Response Evaluation Criteria in Solid Tumors RECIST 1.1 criteria.

**Duration of Approval for Initials and Renewals:** 3 months
**Ceritinib**

**Brand(s): Zykadia**

**DOSAGE FORM/STRENGTH: 150mg capsule**

**Initial Criteria:**

For the treatment of anaplastic lymphoma kinase (“ALK”) – positive locally advanced (not amenable to curative therapy) or metastatic non-small cell lung cancer (NSCLC) in patients meeting ALL the following criteria:

- Patient is using ceritinib as monotherapy; AND
- Ceritinib is being used as second line\(^1\) therapy in patients who have experienced disease progression to crizotinib OR in patients who have experienced intolerance\(^2\) to crizotinib.

**Exclusion criteria:**

- Ceritinib will not be funded if the patient has experienced disease progression while on an ALK inhibitor other than crizotinib.
- Ceritinib will not be funded beyond third line therapy.\(^2\)

\(^1\)Time-limited funding will be considered case-by-case in patients with ALK-positive NSCLC who have progressed on chemotherapy and crizotinib OR crizotinib and an immune checkpoint inhibitor commenced prior to the public funding of ceritinib.

\(^2\)Include details of the intolerance including the grade of toxicity and reasons why crizotinib was not able to be used, particularly in situations where a toxicity was deemed to be grade 1 or 2.

**Recommended dose:** 450 mg daily (Product Monograph dose update December 2018)

**Renewal Criteria:**

Ongoing funding will be considered in patients who have not experienced disease progression or unacceptable toxicities to treatment with ceritinib.

**Approval duration of initial and renewal requests:** 1 year
Cobimetinib
Brand(s): Cotellic
DOSAGE FORM/ STRENGTH: 20 mg tablet

Initial criteria:

For the treatment of patients with previously untreated BRAF V600 mutation-positive unresectable stage III or stage IV melanoma who have a good performance status (ECOG ≤ 2).

- As first-line combination therapy with vemurafenib; AND
- If brain metastases are present, they should be asymptomatic or stable

Recommended Dose as combination dual therapy with Vemurafenib:
Cobimetinib 60 mg once daily for 21 days, followed by seven days off treatment; AND Vemurafenib 960 mg twice daily for 28 days.
Both drugs are given until disease progression or unacceptable toxicity.

Renewal criteria:

Combination dual therapy may be continued until evidence of disease progression\(^1\) or development of unacceptable toxicity requiring discontinuation.

\(^1\) Letter from physician outlining radiological and clinical benefit requiring continuation of the drug and verification of no disease progression or development of unacceptable toxicity must be submitted.

Approval duration (both initial and renewal requests): 6 months (patients should have their disease status assessed at least every 6 months)

Exclusion Criteria:

- BRAF V600 negative, or wild type tumors, or unknown status will not be funded

Cobimetinib therapy will not be considered for funding in patients who have progressed on a prior BRAF inhibitor therapy used as monotherapy or in combination.
Initial Criteria:

For the treatment of locally advanced (not amenable to curative therapy) or metastatic non-small cell lung cancer (NSCLC) in patients who meet the following criteria:

1. Crizotinib is being used as the first-line oral treatment for a patient with ROS1-positive NSCLC

OR

Crizotinib is being used as the first-line oral treatment for a patient with anaplastic lymphoma kinase (ALK)-positive NSCLC

OR

Crizotinib is being used as second line oral treatment for a patient with anaplastic lymphoma kinase (ALK)-positive NSCLC

Note: Patients with ALK-positive NSCL who have progressed on Crizotinib in first line will not be eligible for funding in second line.

2. Patient has good performance status.

3. Patient is using crizotinib as monotherapy.

Renewal Criteria:

Renewals will be considered until clinically meaningful disease progression or the patient has experienced unacceptable toxicity.

Exclusion Criteria:

Crizotinib will not be funded as combination therapy with another treatment for the treatment of ROS1-positive NSCLC or for ALK-positive NSCLC.

Dosing: 250 mg orally twice daily

Approval duration of Initials and Renewals: 1 year
Initial Criteria:

For the mutation-targeted treatment of patients with BRAF V600 mutation-positive unresectable melanoma or metastatic melanoma meeting the following criteria:

- As monotherapy or as combination therapy with trametinib;
- If brain metastases are present, they should be asymptomatic or stable

Exclusion Criteria:

- BRAF V600 negative, or wild type tumors, or unknown status will not be funded
- Funding will not be considered in patients who have experienced progression on a BRAF mutation targeted therapy. The Ministry will fund only one BRAF mutation targeted treatment/treatment regimen.
- May be sequenced after immunotherapies or other funded treatments, however, treatment beyond third line will not be considered for funding.

Recommended Dose as Monotherapy:

150 mg twice daily until disease progression or development of unacceptable toxicity requiring discontinuation of dabrafenib

Recommended Dose as combination dual therapy with Trametinib:

Dabrafenib 150 mg twice daily and Trametinib 2mg once daily until disease progression or development of unacceptable toxicity requiring discontinuation

Renewal Criteria:

Therapy as monotherapy OR as combination dual therapy (as above) may be continued until evidence of disease progression or development of unacceptable toxicity requiring discontinuation.

1 Letter from physician outlining radiological and clinical benefit requiring continuation of the drug and verification of no disease progression must be submitted.

Approval duration (both initial and renewal requests): 6 months (patients should have their disease status assessed at least every 6 months)

Case by case:
Requests in patients who have initiated another single-agent BRAF or MEK inhibitor therapy will be considered on a **case-by-case** basis ONLY IF there has been no disease progression.

**Dabrafenib**  
**Brand(s):** Tafinlar  
**DOSAGE FORM/ STRENGTH:** 50 mg and 75 mg capsule

**For the adjuvant treatment of resected Stage III cutaneous melanoma** in patients meeting ALL the following criteria;

i) Dabrafenib is being used as combination therapy with Trametinib  

ii) Patient’s cutaneous melanoma met the following requirements prior to resection:  
   a. Histologically confirmed stage IIIA (limited to lymph node metastases of > 1 mm), IIIB, IIIC, or IIDD cutaneous melanoma  
   [8th edition of the American Joint Committee on Cancer staging system]  
   b. BRAF V600 mutated (all BRAF V600 mutations)  

iii) Post-resection, clinical or radiographic confirmation of complete disease resection including absence of in-transit metastases must be provided.  

iv) Eastern Cooperative Oncology Group (ECOG) Performance Status 0 to 2  

1Micrometastatic lymph node involvement detected by sentinel lymph node biopsy will be allowed.

**Exclusion Criteria:**
- Patients with stage IIIA cutaneous melanoma with lymph node metastases less than 1 mm.  
- Monotherapy with Dabrafenib  
- Combinations with other anticancer therapies

**Recommended Dose in adjuvant therapy:**  
Dabrafenib 150 mg twice daily and Trametinib 2mg once daily

Notes: Treatment administered post-resection until disease recurrence or unacceptable toxicity to a maximum of 12 months of treatment in total. (Note: 12 months refers to duration of adjuvant treatment accessed through all sources of funding (i.e. private and public).  

**Approval duration:** Maximum of 12 months. Renewals are not considered.

Additional notes:  
1. The Ministry will reimburse for provincially funded treatments for use in the adjuvant setting in patients with cutaneous melanoma for a total duration of up to 12 months. Overall access to adjuvant therapy will be limited to 12 months in total and combines the duration of use of all treatments administered in the adjuvant setting.
2. Funding will not be granted for dabrafenif-trametinib in patients who have used another treatment in the adjuvant setting for cutaneous melanoma for a duration of 3 months or longer.

**Dabrafenib**

**Brand(s):** Tafinlar  
**DOSAGE FORM/ STRENGTH:** 50 mg and 75 mg capsule

3. A one-time switch to dabrafenif-trametinib in the adjuvant cutaneous melanoma setting will be permitted for BRAF-mutated patients who switch from another adjuvant treatment that has been used for less than 3 months and upon meeting the above funding criteria.

4. Patients who experience disease progression within 6 months of completion of adjuvant BRAF therapy will not be funded for another BRAF targeted therapy.

5. Switching to cobimetinib-vemurafenib from dabrafenif-trametinib will not be permitted in BRAF-positive patients.

**Darolutamide**

**Brand(s):** Nubeqa  
**DOSAGE FORM/ STRENGTH:** 300 mg tablet  
**Effective date:** June 8, 2021

**Initiation Criteria:**

For the treatment of high risk non-metastatic castration resistant prostate cancer (nmCRPC) in patients who meet all the following criteria:

1. Patient using darolutamide (Nubeqa) in combination with androgen deprivation therapy (ADT); AND

2. Has no detectable distant metastases as determined by CT, MRI, or technetium-99m bone scan; AND

3. Patient has castrate resistant disease based on meeting all the following indicia observed while on continuous ADT treatment or post orchiectomy:
   - Castrate serum testosterone levels: AND
   - Biochemical progression defined as three (3) prostate-specific antigen (PSA) rises at least 1 week apart, with the last PSA greater than 2ng/mL (if the patient has a history of antiandrogen use, the most recent PSA value must be obtained at least 4 weeks after anti-androgen withdrawal); AND
4. Patient is at high risk for developing metastatic disease based on a prostate-specific antigen doubling time (PSADT) of less than or equal to 10 months during continuous ADT.

5. Has an Eastern Cooperative Oncology Group (ECOG) Performance Status less than or equal to 2.

**Darolutamide**

**Brand(s):** Nubeqa  
**DOSAGE FORM/ STRENGTH:** 300 mg tablet

**Exclusion Criteria:**  
Patients meeting one or more of the below exclusion criteria will not be funded.

- The patient received prior chemotherapy or immunotherapy for the treatment of prostate cancer, unless it was in the adjuvant or neoadjuvant setting completed more than 2 years previously.

- The patient has experienced disease progression on prior treatment with Erleada (apalutamide) or Xtandi (enzalutamide).

- The patient has a high risk for disease progression by other definitions (such as a high Gleason score 8-10, high PSA level at diagnosis, etc.) AND has not had a PSA progression in the non-metastatic setting.

**Approved Dosage:** 600 mg administered orally twice daily.

**Notes:**

- The Ministry will fund only one of Nubeqa (darolutamide) or Erleada (apalutamide) or Xtandi (enzalutamide) in patients with non-metastatic castrate resistant prostate cancer.

- While doses may be withheld or reduced to 300 mg twice daily to manage grade 3 toxicities until symptoms improve, treatment doses should be resumed at a dose of 600 mg twice daily.

- Requests for Nubeqa in patients who initiated Erleada or Xtandi therapy in the nmCRPC setting and who have not had disease progression will be considered on a case-by-case basis.

- Following progression on darolutamide in non-metastatic castrate resistant prostate cancer, the patient would not be eligible for darolutamide, apalutamide or enzalutamide in metastatic castrate resistant state.

**Renewal Criteria:**
Renewals will be considered in patients without evidence of radiographic disease progression or unacceptable toxicity while on Nubeqa therapy.

**Approved Dosage:** 600 mg administered orally twice daily.

**Approval Duration of initials and renewals:** 1 year
Dasatinib
Brand(s): Sprycel and generics (see formulary for list of funded generics)
DOSAGE FORM/ STRENGTH: 20 mg, 50 mg, 70 mg, 100 mg tablet

For the treatment of Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML) in the chronic phase.

Dosing recommendation: 100 mg per day.

Renewals will be considered for patients who have experienced hematologic and/or cytogenic response and is expected to continue to do so.

Duration of Approval: 1 Year

Exclusion criteria:

Combination treatment with any two or more of the oral tyrosine-kinase inhibitors (TKI) (i.e. imatinib, nilotinib or dasatinib) will not be funded.

Note: Funding is only considered for any two oral TKIs* per patient in a lifetime for chronic phase CML (*TKIs: imatinib, nilotinib, or dasatinib). If a patient develops grade 3 or grade 4 toxicity on one of the listed TKI’s within 3 months of initiating therapy, funding for a third oral TKI will be allowed.

For the treatment of patients with accelerated phase or blast phase Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML) with documented resistance or intolerance (as defined below) to imatinib therapy

Dosing recommendation: 140 mg per day.

Definitions of resistance and intolerance:

Imatinib resistance is defined as primary or acquired resistance to imatinib at doses of at least 600 mg/day or through a mutational analysis report.

Intolerance to imatinib (at any dose) is defined as persistent grade 3 or grade 4 toxicity requiring discontinuation of therapy.

Renewals will be considered for patients who have experienced hematologic and/or cytogenic response and are expected to continue to do so.

Duration of Approval: 1 Year
Dasatinib
Brand(s): Sprycel and generics (see formulary for list of funded generics)
DOSAGE FORM/STRENGTH: 20 mg, 50 mg, 70 mg, 100 mg tablet

Exclusion criteria:

- Combination treatment with any 2 or more of the oral TKIs (i.e. imatinib, nilotinib or dasatinib) will not be funded.
- Dasatinib is not funded as a sequential third line therapy in patients who experience primary or acquired resistance (not including mutational resistance) to nilotinib.

For the treatment of Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ ALL) in patients meeting the following criteria:

i) An adult patient with Philadelphia chromosome positive acute lymphoblastic leukemia (Ph +ALL); AND

ii) Patient’s disease is resistant\(^1\) to imatinib-containing chemotherapy (patient must have tried 600 mg/day); OR

iii) Patient has experienced intolerance\(^2\) to imatinib therapy.

\(^1\)Imatinib resistance is defined as primary or acquired resistance to imatinib at doses of at least 600 mg/day or through a mutational analysis report.

\(^2\)Intolerance to imatinib (at any dose) is defined as the patient has experienced persistent grade 3 or grade 4 toxicity requiring discontinuation of therapy.

Renewals will be considered after confirmation from the patient’s physician that the patient has benefited or continues to benefit from therapy with Sprycel and is expected to continue to do so.

Duration of Approval: 1 Year

Reimbursement of dasatinib for children with acute lymphoblastic leukemia will be considered on a case-by-case basis.
Entrectinib
Brand(s): Rozlytrek
DOSAGE FORM/STRENGTH: 100 mg, 200 mg capsule
Effective date: December 23, 2021

Initial Criteria:

For the treatment of locally advanced (not amenable to curative therapy) or metastatic non-small cell lung cancer (NSCLC) in patients who meet the following criteria:

1. Entrectinib is being used as the first-line oral treatment for a patient with ROS1-positive NSCLC; AND
2. Patient has good performance status; AND
3. Patient is using entrectinib as monotherapy.

1Include the ROS1 laboratory report with the application.

Renewal Criteria:

Renewals will be considered until clinically meaningful disease progression or the patient has experienced unacceptable toxicity.

Exclusion Criteria:

1. Entrectinib will not be funded as combination therapy with another treatment for the treatment of ROS1-positive NSCLC.
2. Entrectinib will not be funded in patients who have previously experienced disease progression while on crizotinib for ROS1-positive NSCLC.

Notes:

1. Patients who have initiated on a first-line platinum-based doublet chemotherapy, chemotherapy-immunotherapy combination, or single agent immunotherapy and have not progressed, may be considered for entrectinib if they have not received an ROS-1 targeted treatment previously.

Approved Dose: Maximum of 600 mg orally once daily

Approval duration of Initials and Renewals: 1 year
Initiation criteria for metastatic castration sensitive prostate cancer (mCSPC):

For the treatment metastatic castration sensitive prostate cancer (mCSPC) in patients who meet all of the following criteria:

1. Enzalutamide is used in combination with androgen deprivation therapy (ADT)\(^1\); AND
2. Metastatic lesions detected on technetium-99m bone scan, computed tomography (CT), and/or magnetic resonance imaging (MRI); AND
3. Castration sensitive as defined by the patient being treatment naïve to an ADT OR ADT initiated within the prior 6 months before start of therapy with enzalutamide OR ADT used in the neoadjuvant or adjuvant setting where the patient has been off treatment for 12 months or more prior to start of enzalutamide; AND
4. Has not experienced disease progression with another an androgen receptor axis targeted therapy (ARAT) for castration sensitive prostate cancer; AND
5. Patient has good performance status.

\(^1\)ADT is not required for patients with bilateral orchiectomy

Definitions for the purpose of the EAP funding criteria:

**ADT** – A first generation androgen deprivation therapy (e.g. goserelin, leuprolide, triptorelin, buserelin, degarelix)

**ARI**- A second generation androgen receptor inhibitor (e.g. apalutamide, darolutamide, enzalutamide)

**ARAT**- An androgen receptor axis targeted therapy (e.g. abiraterone, apalutamide, darolutamide, enzalutamide)

The following baseline levels are to be provided with the initial application;

i) Number of metastatic lesions on bone scan and in soft tissues

ii) Testosterone level

iii) Baseline (pre-treatment) PSA level

iv) Pre-treatment Gleason score (optional)
Enzalutamide
Brand(s): Xtandi
DOSAGE FORM/STRENGTH: 40 mg capsule

Notes:

1. Patients who have previously progressed on a second-generation androgen receptor inhibitor (e.g. apalutamide, enzalutamide, darolutamide) for prostate cancer will not be eligible for enzalutamide in mCSPC. The Ministry will fund only one of enzalutamide or apalutamide in patients with mCSPC.

2. Patients treated with apalutamide or darolutamide as part of a clinical trial may be eligible for enzalutamide and will be considered on a case-by-case basis.

3. Patients who progress on treatment with enzalutamide or apalutamide mCSPC will not be eligible for enzalutamide in the metastatic castration resistant setting.

4. Patients who are currently on a treatment regimen with an androgen receptor axis targeted therapy (e.g. abiraterone or a second-generation androgen receptor inhibitor) for prostate cancer, must meet the initiation criteria if they wish to switch to publicly funded enzalutamide for mCSPC.

5. Time limited funding consideration will be provided on a case-by-case basis for those patients who are using docetaxel in combination with ADT as long as there has been no progression and the treatment regimen has not been used for more than 6 months.

Approved Dosage: 160 mg administered orally once daily.

Renewal Criteria:

Renewals will be considered in patients until disease progression (i.e. PSA elevation in addition to radiographic disease progression, or PSA progression in addition to clinical symptoms associated with cancer progression), development of castration resistant disease, or experiencing unacceptable toxicity while on enzalutamide.

Exclusion Criteria: (Patients meeting any of the following will not be funded.)

1. Patients who have previously experienced disease progression on enzalutamide or another ARI used in the setting of prostate cancer.

2. Patients who have risk factors for seizures

3. Enzalutamide will not be funded as combination therapy with another androgen receptor axis targeted therapy.

Approved Dosage for Initials and Renewals: 160 mg administered orally once daily.

Duration of initial and renewal approvals: 1 year
Initiation Criteria for high risk non-metastatic castration resistant prostate cancer (nmCRPC):

For the treatment of high risk non-metastatic castration resistant prostate cancer (nmCRPC) in patients who meet all the following criteria:

1. Patient using Xtandi in combination with androgen deprivation therapy (ADT); AND

2. Has no detectable distant metastases as determined by CT, MRI, or technetium-99m bone scan; AND

3. Patient has castration resistant disease based on meeting all the following indicia observed while on continuous ADT treatment or post orchiectomy:
   - Castrate serum testosterone levels: AND
   - Biochemical progression defined as Three (3) prostate-specific antigen (PSA) rises at least 1 week apart, with the last PSA greater than 2ng/mL; and

4. Patient is at high risk for developing metastatic disease based on a Prostate-specific antigen doubling time (PSADT) of less than or equal to 10 months during continuous ADT.

5. Has an Eastern Cooperative Oncology Group (ECOG) Performance Status less than or equal to 2.

Approval Duration: 1 year

Exclusion Criteria:

1. The patient received prior chemotherapy for the treatment of prostate cancer, unless it was in the adjuvant or neoadjuvant setting.

2. The patient has previously experienced disease progression on apalutamide or darolutamide used for prostate cancer.

3. The patient has risk factors for seizures.
Enzalutamide
Brand(s): Xtandi
DOSAGE FORM/STRENGTH: 40 mg capsule

Definitions for the purpose of the EAP funding criteria:

**ADT** – A first generation androgen deprivation therapy (e.g. goserelin, leuprolide, triptorelin, buserelin, degarelix)

**ARI**- A second generation androgen receptor inhibitor (e.g. apalutamide, darolutamide, enzalutamide)

**Approved Dosage**: 160mg administered orally once daily

**Notes:**

1. The Ministry will fund only one second generation androgen receptor inhibitor (e.g. apalutamide, darolutamide, or enzalutamide) in patients with nmCRPC.

2. Patients who have progressed on enzalutamide in nmCRPC will not be eligible for enzalutamide in metastatic castration resistant prostate cancer (mCRPC).

3. Requests for enzalutamide in patients who have initiated another ARI therapy in the nmCRPC setting and who have not experienced disease progression will be considered on a case-by-case basis.

**Renewal Criteria:**

Renewals will be considered in patients without evidence of radiographic disease progression or unacceptable toxicity while on enzalutamide therapy.

**Approval Duration**: 1 year
Initiation criteria for metastatic castration resistant prostate cancer (mCRPC):

For the treatment of metastatic castration resistant prostate cancer (mCRPC) in patients meeting the following criteria:

1. Enzalutamide is used in combination with androgen deprivation therapy (ADT); AND
2. Metastatic lesions detected on a bone scan, computed tomography (CT), and/or magnetic resonance imaging (MRI); AND
3. Patient has castration resistant disease based on meeting the following indicia observed while on continuous ADT treatment or post orchiectomy:
4. Castrate serum testosterone levels; AND
5. Biochemical progression defined as three (3) prostate-specific antigen (PSA) rises at least 1 week apart, with the last PSA greater than 2ng/mL AND/OR radiographic progression of new or pre-existing disease as determined by the detection of 2 or more lesions on bone scan or presence of new soft tissue lesions by RECIST criteria; AND
6. Patient has progressed on a docetaxel-based chemotherapy
   OR
   Patient is using enzalutamide pre-docetaxel for metastatic castration resistant prostate cancer and has not previously experienced disease progression on enzalutamide or another second generation androgen receptor inhibitor (e.g. apalutamide, darolutamide,) used for prostate cancer
   OR
   Patient is using enzalutamide pre-docetaxel for mCRPC and has not previously experienced disease progression on abiraterone used in mCRPC sequenced immediately prior to enzalutamide;

   AND
7. Patient has good performance status defined as an Eastern Cooperative Oncology Group (ECOG) Performance status less than or equal to 2.

1ADT is not required for patients with bilateral orchiectomy

Definitions for the purpose of the EAP funding criteria:

**ADT** – A first generation androgen deprivation therapy (e.g. goserelin, leuprolide, triptorelin, buserelin, degarelix)

**ARI** - A second generation androgen receptor inhibitor (e.g. apalutamide, darolutamide, enzalutamide)

**ARAT** - An androgen receptor axis targeted therapy (e.g. abiraterone, apalutamide, darolutamide, enzalutamide)
Enzalutamide
Brand(s): Xtandi
DOSAGE FORM/STRENGTH: 40 mg capsule

Notes:

1. Patients who have progressed on treatment with second generation androgen receptor inhibitors (e.g. apalutamide, enzalutamide, darolutamide) used in prior stages of prostate cancer will not be eligible for enzalutamide in the metastatic castration resistant setting.
2. Patients treated with apalutamide or darolutamide as part of a clinical trial may be eligible for enzalutamide and will be considered on a case-by-case basis.
3. Requests for enzalutamide in patients who initiated abiraterone therapy and who have not had disease progression while on abiraterone will be considered on a case-by-case basis.

Exclusion criteria: (Patients meeting any of the following will not be funded)

1. The patient has risk factors for seizures;
2. Enzalutamide will not be funded as combination therapy with another androgen receptor axis targeted therapy or cabazitaxel.

Renewals:
Renewals will be considered in patients who have not experienced disease progression while on enzalutamide therapy.

Approved Dosage for Initials and Renewals: 160 mg administered orally once daily.

Duration of initial and renewal approvals: 1 year
Erlotinib
Brand(s): Tarceva and generics see formulary for a funded list of generics
DOSAGE FORM/ STRENGTH: 25 mg, 100 mg, 150 mg tablet

For the treatment of clinically documented incurable progressive non-small cell lung cancer (NSCLC) where:

- Erlotinib is used as monotherapy for the 2nd- or 3rd-line treatment after failure of prior chemotherapy (any regimen) in patients 70 years of age or older.

- Erlotinib is used as monotherapy for the 2nd- or 3rd-line treatment of patients with clinically documented incurable progressive non-small cell lung cancer (NSCLC) despite prior chemotherapy including both docetaxel and a platinum-based treatment (i.e. cisplatin or carboplatin).

- Erlotinib is used as monotherapy for the 3rd-line treatment of patients with clinically documented incurable progressive non-small cell lung cancer (NSCLC) despite prior chemotherapy including both a platinum-based therapy (i.e. cisplatin or carboplatin) AND either pemetrexed or topotecan.

- Erlotinib is used as monotherapy for 2nd line treatment of NSCLC after 1st line platinum-based therapy, where no other chemotherapy will be given and erlotinib is used as the last treatment for the patient

Patients should be assessed for disease status at least every two months. Erlotinib should be discontinued if there is evidence of disease progression.

Note that erlotinib is not indicated and therefore, is not considered for reimbursement as 1st line therapy in treatment of NSCLC.

Requests for 2nd-line and 3rd-line use of erlotinib in patients 70 years of age or older and have not received treatment with either platinum-based combinations will be considered on a case-by-case basis.

Approved dosage: 150 mg/day

Duration of Approval: 6 Months

Renewal will be considered for patients who respond to therapy with no evidence of disease progression. Patients should be assessed for disease status at least every two months. Erlotinib should be discontinued if there is evidence of disease progression.

Duration of Approval: 6 Months
**Everolimus**  
Brand(s): Afinitor and generics see formulary for a funded list of generics  
DOSAGE FORM/ STRENGTH: 2.5 mg, 5 mg, 10 mg tablet

For the treatment of metastatic renal cell carcinoma (mRCC) as second or third line\(^1\) therapy in patients previously treated for mRCC with a funded tyrosine kinase inhibitor (TKI).

Exclusion criteria: Use in the 4\(^{th}\) line setting or later in the treatment course of their disease  
Dosage: 10 mg daily

**Renewal** will be considered for those who have demonstrated benefit from Afinitor therapy (i.e. no disease progression) and is expected to continue to do so.

\(^1\)Funded TKIs include sunitinib (Sutent), sorafenib (Nexavar), and pazopanib (Votrient). The criteria are derived from the review of everolimus for provincial funding for the treatment of MRCC at the time of the original review. Drugs that may have been used as standard treatment in first line may have included interferon and temserolimus. Everolimus is currently not funded after progression on axitinib (Inlyta) or nivolumab.

For the treatment of patients who have progressive, unresectable, well or moderately differentiated, locally advanced or metastatic pancreatic neuroendocrine tumors (pNET).

Patient must have an ECOG\(^*\) ≤ 2 (prior to the start of Afinitor therapy).

*ECOG = Eastern Cooperative Oncology Group Status

Exclusion criteria: the patient’s disease progressed while taking sunitinib (Sutent) to treat pNET.  
Dosage: 10 mg daily

**Duration of Approval:** 1 year

**Renewal** will be considered for those who have benefited from Afinitor therapy (i.e. no disease progression) and is expected to continue to do so.

Reimbursement of Afinitor will be considered until disease progression occurs on Afinitor.  
**Duration of Approval:** 1 year
Everolimus
Brand(s): Afinitor and generics (see formulary for a funded list of generics)
DOSAGE FORM/ STRENGTH: 2.5 mg, 5 mg, 10 mg tablet

For the treatment of unresectable, locally advanced or metastatic, well-differentiated non-functional neuroendocrine tumours (NETs) of gastrointestinal or lung origin (GIL) in adult patients meeting the following criteria:

- Documented radiological disease progression within six months; AND
- Good performance status (ECOG 0-2).

Treatment should continue until confirmed disease progression or unacceptable toxicity.

Renewals will be considered where the patient’s physician has confirmed that the Patient has benefited or continues to benefit from therapy with Afinitor as evidenced by no disease progression, and that they are expected to continue to do so.

For the treatment of postmenopausal women with hormone-receptor positive, HER2 negative advanced breast cancer meeting the following criteria:

- Afinitor is to be used in combination with exemestane; AND
- Patient must have an ECOG* ≤ 2 after recurrence or progression following a non-steroidal aromatase inhibitor (NSAI).

  *ECOG = Eastern Cooperative Oncology Group Status

Dosage: 10 mg daily (dose titration is allowed).

**Duration of Approval:** 1 year

**Renewals** will be considered for patients who have benefited or continues to benefit from therapy with Afinitor and is expected to continue to do so.

**Duration of Approval:** 1 year
Everolimus  
Brand(s): Afinitor and generics (see formulary for a funded list of generics)  
DOSAGE FORM/ STRENGTH: 2.5 mg, 5 mg, 10 mg tablet  

For the treatment of renal angiomyolipoma (AML) associated with tuberous sclerosis complex (TSC) in patients who meet all the following conditions:

(i) Presence of coalescent or multifocal AMLs in either one or both kidneys; AND

(ii) AML progression despite previous embolization and/or surgery; AND

(iii) Further embolization and/or surgery is not recommended due to a documented clinical reason (Note: The physician must submit a clinical note with the request outlining/detailing why invasive therapy cannot be considered);

The approved dosage: 10 mg orally once daily.

Duration of Approval: 1 year

Case-by-Case consideration will be considered in patients who have never been treated with invasive procedures such as embolization and/or surgery. The physician must provide detailed clinical rationale (e.g., from clinical consultation notes) as to why embolization and/or nephrectomy would be medically contraindicated for the patient.

Renewals will be considered in patients with the following documented benefits from therapy;

No AML progression (i.e. no significant new lesions and increase in kidney volume, as well as no significant AML related bleeding);

AND
  • There is a reduction in volume of AMLs identified prior to treatment with the everolimus.

Duration of Approval: 2 years
Everolimus
Brand(s): Afinitor and generics (see formulary for a funded list of generics)
DOSAGE FORM/ STRENGTH: 2.5 mg, 5 mg, 10 mg tablet

For the treatment of Subependymal giant cell astrocytoma (SEGA) associated with tuberous sclerosis complex (TSC) for whom surgical resection cannot be considered* for reasons such as:
- Location, size, and/or distribution of tumour(s); OR
- SEGA progression despite previous surgical interventions; OR
- Neurocognitive problems/ other complications secondary to previous surgical interventions.
- *Requests must provide details/ consultation notes outlining why the patient cannot be considered for surgical treatment.

Duration of Approval: 1 year

Renewals will be considered in patients with the following documented benefits from therapy:
- Stabilization of SEGA progression (based on assessment of SEGA volume and/or appearance of new lesions); AND
- Improvement of symptoms (e.g., reduced seizure frequency and decreased need for neurosurgical intervention).

Duration of Approval: 2 years
**Gefitinib**

**Brand(s):** Iressa and generics (see formulary for funded OFIs)

**DOSAGE FORM/ STRENGTH:** 250 mg tablet

For the first line, monotherapy treatment of locally advanced (not amenable to curative therapy) or metastatic non-small cell lung cancer (NSCLC) in patients who have activating mutations of epidermal growth factor receptor-tyrosine kinase (EGFR-TK). (i.e. Patients who are EGFR Positive)

The patient is to be assessed for disease status at least every two months and treatment will be discontinued if there is evidence of disease progression.

Dose Reimbursed: 250 mg orally once daily.

**Duration of Approval:** 6 months

Iressa will not be granted funding in the following circumstances;

Patients with EGFR wild-type mutation (i.e. negative for mutation);

Patients with EGFR unknown mutation;

2\(^{nd}\) or 3\(^{rd}\) line or maintenance NSCLC; or

Patients with unknown EGFR status who start their first chemotherapy while waiting for EGFR testing, then are found/confirmed to be EGFR positive, should continue with the current therapy and will not be eligible for gefitinib (Iressa) in this setting.

Patients who receive gefitinib (Iressa) first line are not eligible for erlotinib in the second- or third-line in the setting of maintenance therapy of NSCLC.

Requests for gefitinib for patients who have initiated another EGFR TKI therapy (i.e. Afatanib [Giotrif]) in the first line setting and who have not had disease progression will be considered on a case-by-case basis.

**Renewal** will be considered for patients until there is any evidence of disease progression, at which point, treatment with gefitinib (Iressa) must be discontinued. Patients must have their disease status assessed at least every two months.

Dose Reimbursed: 250 mg orally once daily.

**Duration of Approval:** 6 months
Gilteritinib
Brand(s): Xospata
DOSAGE FORM/ STRENGTH: 40 mg tablet
Effective date: December 1, 2021

For the treatment of adult patients diagnosed with relapsed or refractory FMS-like tyrosine kinase 3 (FLT3)-mutated acute myeloid leukemia (AML) who meet the following criteria;

1. Patient has relapsed or is considered to be refractory to a prior chemotherapy regimen used for AML.
2. FLT3 mutation with either a FLT3-ITD, FLT3-TKD/D835 or FLT3-TKD/I836 is confirmed by an approved test taken after relapse on a chemotherapy regimen; AND
3. Patient has good performance status (e.g. ECOG less than or equal to 2).

Exclusion criteria: (Requests meeting ANY of the following criteria will not be funded.)

1. Patients with therapy-related AML.
2. Patients who have had a prior relapse while being treated with gilteritinib or another tyrosine kinase inhibitor (TKI) specific to FLT3-mutated AML in the relapsed or refractory AML setting.
3. Gilteritinib is not funded for induction or consolidation therapy for AML (i.e. not funded for earlier lines of treatment prior to refractory or relapsed disease).

Renewal Criteria:

Treatment to continue until disease progression or until unacceptable toxicity occurs. At the time of renewal please include a brief summary of the clinical benefit that has been observed (e.g. partial or complete remission, reduction in transfusions etc.)

Notes:

1. A delay in clinical response can occur and it is recommended that treatment with gilteritinib be continued for a minimum of six months (i.e. in the absence of disease progression or unacceptable toxicity).
2. Patients may be considered to have refractory AML after progression on a minimum of one cycle of induction chemotherapy in accordance with the prescriber’s clinical assessment.
3. Dose escalations to achieve complete remission will be permitted but dose should not be escalated after achieving complete remission.
4. Patients who receive gilteritinib in relapsed or refractory AML who progress to a hematopoietic stem cell transplant will be able to continue on gilteritinib as maintenance until disease progression.
For a time-limited period, patients currently receiving salvage chemotherapy for relapsed or refractory AML and patients in second hematologic relapse or later may be considered for funding on a case-by-case basis if they did not relapse on a prior TKI in the relapse/refractory setting if they have FLT3 mutated AML.

Recommended dose: 120mg once daily

Duration of approval of initial requests: 7 months

Duration of approval or renewals: 1 year
Ibrutinib
Brand(s): Imbruvica
DOSAGE FORM/ STRENGTH: 140 mg capsule

For the treatment of patients with chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL) who meet the following criteria:

i) Patient has received at least one prior therapy to treat CLL/SLL; AND

ii) Patient’s prescriber has deemed that it would be inappropriate for the patient to receive treatment or retreatment with a fludarabine-based regimen.

Duration of Approval: 1 Year

Exclusion criteria:

Patients whose disease has progressed on idelalisib therapy in the relapsed setting are not eligible to receive iburitinib.

Renewals will be considered for patients who have not experienced disease progression while on ibrutinib (Imbruvica) therapy.

Duration of Approval: 1 Year

Initial criteria for Treatment naïve patients with high risk CLL/SLL (First-line therapy):

For patients with previously untreated chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL) who present with one of the following cytogenic markers:

- chromosome 17p deletion; OR
- TP 53 mutation; OR
- unmutated immunoglobulin heavy chain variable region (IgHV)

Renewal criteria: Patient has experienced no disease progression while on Imbruvica therapy.

Initial and renewal approval period: 1 year.

For treatment of patients with relapsed or refractory mantle cell lymphoma who have received at least one prior therapy.

Renewals will be considered if patient has experienced no disease progression while on Imbruvica therapy.

Initial and renewal approval period: 1 year.
**Idelalisib**

**Brand(s):** Zydelig  
**DOSAGE FORM/ STRENGTH:** 100 mg, 150 mg Tablets

For the treatment of patients with relapsed chronic lymphocytic leukemia (CLL)/small lymphocytic lymphoma (SLL) in combination with Rituximab.

Exclusion criteria:

Patients whose disease has progressed on ibrutinib therapy in the relapsed setting are not eligible to receive idelalisib.

Note: Patients who have experienced intolerance but not disease progression to ibrutinib in the relapsed setting may switch to idelalisib. Documentation on the nature of the intolerance is required.

**Renewals** will be considered for patient who has not experienced disease progression while on idelalisib (Zydelig) therapy.

**Funded Dose:**

Idelalisib will be funded in combination with up to 8 cycles of rituximab at the recommended dose of 150 mg orally twice daily and will continue following the completion of the rituximab portion of the regimen.

**Imatinib**

**Brand(s):** Gleevec + generics (see below for billing information)  
**DOSAGE FORM/ STRENGTH:** 100 mg tablet, 400 mg tablet

For the treatment of Metastatic Gastrointestinal Stromal Tumours (GIST) in patients with a tumour deemed to be NOT surgically resectable (metastatic or recurrent)

**Duration of Approval:** 1 Year

**Renewal** will be considered for patients with GIST who have benefited from or continues to benefit from therapy with Gleevec and is expected to continue to do so.

**Duration of Approval:** 1 Year

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<th>Drug Product</th>
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Imatinib
Brand(s): Gleevec + generics (see below for billing information)
DOSAGE FORM/ STRENGTH: 100 mg tablet, 400 mg tablet

For the Adjuvant treatment of Gastrointestinal Stromal Tumours (GIST) in patients who meet the following criteria:

Patients are at intermediate to high risk of recurrence following complete resection (using Miettinen relapse risk criteria, risk ≥ 20%) or has had tumor rupture before surgery or at surgery; AND

- The pathology has been confirmed with c-kit positivity.

Note that the dosing regimen covered is no more than 400 mg daily.

Duration of Approval: 3 Years

Renewals will NOT be considered for patients receiving Gleevec for Adjuvant GIST.

(i.e. Funding for adjuvant GIST is approved for up to 3 years. Longer coverage durations are not considered.)

For the treatment of adult patients with newly diagnosed Philadelphia chromosome-positive (Ph+) acute lymphoblastic leukemia (ALL)

Duration of Approval: 1 Year

Renewal will be considered for patients receiving Gleevec for Philadelphia chromosome-positive (Ph+) acute lymphoblastic leukemia (ALL) who demonstrate a hematologic or cytogenetic response to therapy

Duration of Approval: 1 Year

As of June 15, 2013, EAP approval letters will indicate PINs to be used for billing purposes. The PINs will allow the full price of each product to be submitted for reimbursement of EAP approved requests. Pharmacists should refer to the respective product monograph(s) for prescribing information and approved indications.

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<th>Drug Product</th>
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Lapatinib
Brand(s): Tykerb
DOSAGE FORM/ STRENGTH: 250 mg tablet
Effective date: November 14, 2011

For the second-line treatment of HER2-positive metastatic breast cancer when used in combination with chemotherapy after previous exposure to trastuzumab-based treatments.

For the treatment of HER-2 positive metastatic breast cancer when used in combination with chemotherapy after use of trastuzumab in patients who have an adverse drug reaction or contraindication to trastuzumab therapy.

Lapatinib will not be considered in patients who meet the following exclusions:

- Lapatinib (Tykerb) will not be funded in combination with trastuzumab (Herceptin) for second-line HER-2 positive metastatic breast cancer.

- Patients who have progressed while on trastuzumab (Herceptin) for second-line treatment of HER-2 positive metastatic breast cancer, will not be eligible for funding of lapatinib (Tykerb)

- Lapatinib (Tykerb) will not be funded in the adjuvant setting.

Dosing schedule:

- 1250 mg (5 tablets) once daily in combination with capecitabine for days 1 to 14 (in a 21 day cycle) until disease progression, unacceptable toxicity, or withdrawal of consent

Note: Funding of second-line lapatinib for HER-2 positive metastatic breast cancer will be discontinued upon evidence of disease progression

Duration of Approval: 6 Months

Renewal will be considered for lapatinib until there is evidence of disease progression at which point the drug should be discontinued.

Duration of Approval: 6 Months
Lenalidomide
Brand(s): Revlimid and generics (see formulary for funded OFIs)
DOSAGE FORM/ STRENGTH: 2.5mg, 5 mg, 10 mg, 15 mg, 20 mg, 25 mg capsule
Updated: December 16, 2020

For the treatment of anemia due to myelodysplastic syndrome (MDS) for patients who meet all the following clinical criteria:

- Demonstrated diagnosis of MDS on bone marrow aspiration; AND
- Presence of del[5q] documented by standard cytogenetic or fluorescence in situ hybridization; AND
- International Prognostic Scoring System (IPSS) risk category low or intermediate-1; AND
- Transfusion-dependent symptomatic anemia.

Duration of Approval: 6 Months

Renewal will be considered for patients who are transfusion-dependent and who have demonstrated at least a fifty percent (50%) reduction in transfusion requirements.

Duration of Approval: Up to 1 Year

Note: Patients with anemia due to MDS who are not transfusion-dependent will be assessed on a case-by-case basis.

- Physicians submitting initial requests for non-transfusion-dependent patients must provide clinical evidence of symptomatic anemia affecting the patient’s quality of life and the rationale for why transfusions are not being used.
- Renewal requests for non-transfusion-dependent patients will be considered on a case-by-case basis. In such cases, the requesting physician will be required to provide the patient’s serial clinical blood culture (CBC) pre- and post-lenalidomide therapy in addition to any other objective evidence of the patient’s response to lenalidomide therapy.
**Lenalidomide**

Brand(s): Revlimid and generics (see formulary for funded OFIs)

DOSAGE FORM/STRENGTH: 2.5mg, 5 mg, 10 mg, 15 mg, 20 mg, 25 mg capsule

For the treatment of patients with multiple myeloma who are not eligible to receive an autologous stem cell transplant who meet all the following criteria;

1. Patient is deemed to be lenalidomide sensitive¹; AND

2. Has good performance status; AND

3. Lenalidomide¹ is being used in one of the following situations;

   i) As first line treatment in previously untreated multiple myeloma within a dual regimen in combination with dexamethasone; OR

   ii) As first line treatment in previously untreated multiple myeloma as part of a triplet regimen in combination with bortezomib² and dexamethasone; OR

   iii) As second line or third line treatment in a patient with relapsed or refractory multiple myeloma in combination with dexamethasone; OR

   iv) As second line or third line treatment in a patient with relapsed or refractory multiple myeloma in combination with dexamethasone and carfilzomib or daratumumab.² Note: Only one novel triplet regimen will be funded for relapsed/refractory disease.

¹Lenalidomide sensitive" is defined as a patient whose disease has not been refractory to a lenalidomide-based regimen and/or has not experienced disease progression while on a lenalidomide-based regimen.

Refractory disease is defined as;

i) disease progression within 60 days after stopping treatment while on lenalidomide (and/or bortezomib); or

ii) progression while on any dose of lenalidomide (and/or bortezomib) including while on maintenance therapy with these therapies; or

iii) non-responsive disease during therapy (either failure to achieve minimal response or experiencing disease progression)

iv) Patients who are refractory to lenalidomide will not be eligible for daratumumab or carfilzomib-based triplets that are used in combination with lenalidomide.
Lenalidomide
Brand(s): Revlimid and generics (see formulary for funded OFIs)
DOSAGE FORM/ STRENGTH: 2.5mg, 5 mg, 10mg, 15mg, 20mg, 25mg capsule

Progressive disease is defined as having one or more of the following:

i) An increase of 25% from lowest response value in serum M-component (the absolute increase must be greater than or equal to 0.5g/dL), and/or urine M-component (the absolute increase must be greater than or equal to 200 mg/24 hours).

ii) An absolute increase of greater than 10mg/dL in the difference between involved and uninvolved FLC levels (if no measurable serum and urine M-protein levels).

iii) Definite development of new bone lesions or soft tissue plasmacytomas or definite increase in the size of existing bone lesions or soft tissue plasmacytomas.

iv) Development of hypercalcemia (corrected serum calcium greater than 11.5 mg/dL or 2.5 mmol/L) that can be attributed solely to the plasma cell proliferative disorder.

2 Patients with multiple myeloma who experience disease progression on a lenalidomide-bortezomib-dexamethasone triplet will not qualify for further lenalidomide-based or bortezomib-based regimens subsequent to disease progression, in both maintenance or relapsed/refractory multiple myeloma.

3 Patients must meet the eligibility requirements for bortezomib, carfilzomib and daratumumab through the New Drug Funding Program (NDFP). Please refer to the Ontario Health – Cancer Care Ontario website for full and update information for funding of NDFP drugs used in for the treatment of multiple myeloma.

Renewal criteria:
Renewals will be considered for lenalidomide for the treatment of multiple myeloma in those who continue to respond to therapy and have not experienced refractory disease or progressive disease while on the lenalidomide-based regimen.

Exclusion Criteria:
Patients meeting the following are not considered for funding:

1. Patients with multiple myeloma who have experienced disease that has been refractory to treatment with a lenalidomide-based treatment.
2. Patients with multiple myeloma who have experienced disease progression while on a lenalidomide-based treatment used for multiple myeloma in any setting including in maintenance treatment.
3. Patients requesting lenalidomide to be used within any regimen who have progressed on 3 lines of treatment with any treatment regimen.
4. Patients with monoclonal gammopathy of uncertain significance (MGUS), smoldering myeloma, or primary amyloidosis.
Lenalidomide
Brand(s): Revlimid and generics (see formulary for funded OFIs)
DOSAGE FORM/STRENGTH: 2.5mg, 5 mg, 10 mg, 15 mg, 20 mg, 25 mg capsule

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

Recommended Dosage:

Initial dose of 10 mg daily. Dose adjustments of 5 mg to 15 mg may be necessary based on individual patient characteristics and responses to lenalidomide.

Duration of Approval: 1 Year

Renewals will be considered until evidence of disease progression or development of unacceptable toxicity to lenalidomide requiring discontinuation of therapy.

Duration of Approval: 1 Year

Additional Notes:

Lenalidomide combinations funded with an injectable anticancer drug through the New Drug Funding Program (NDFP).

Please refer to the CCO website for the full funding criteria for Bortezomib, Carfilzomib and/or Daratumumab triplet combinations with lenalidomide and dexamethasone.

1. Lenalidomide-bortezomib-dexamethasone:

- Bortezomib 1.3 mg/m² or 1.5 mg/m² intravenously (IV) or subcutaneously (SC) days 1, 8, and 15, in combination with lenalidomide and dexamethasone, every 3 weeks for 8 cycles.
- Bortezomib can also be given at 1.3 mg/m² IV or SC on days 1, 4, 8, and 11 every 3 weeks.
- All cycles of bortezomib are given with lenalidomide and dexamethasone. Starting with cycle 9 onwards, lenalidomide and dexamethasone should be continued as maintenance until disease progression or unacceptable toxicity.

Notes:
Regardless of the chosen administration schedule (i.e. once weekly or twice weekly), bortezomib will be funded for a total of eight 3-week cycles.
Patients whose disease is refractory* to both lenalidomide and bortezomib will not be eligible for daratumumab or carfilzomib-based triplets in the relapsed/refractory setting.
**Lenalidomide**

**Brand(s):** Revlimid and generics (see formulary for funded OFIs)

**DOSAGE FORM/ STRENGTH:** 2.5mg, 5 mg, 10 mg, 15 mg, 20 mg, 25 mg capsule

2. **Lenalidomide within a triplet combination with Carfilzomib + Dexamethasone**

Note: The Ministry will fund one novel triplet therapy for relapsed multiple myeloma. Patients who experience toxicity to one of the funded novel triplets may switch once to another triplet within the first 3 months of starting treatment.

**Eligibility Criteria:**
- Carfilzomib is used in combination with lenalidomide and dexamethasone for patients with multiple myeloma who have received at least one prior treatment.
- Treatment should be in patients who have good performance status and are deemed to have adequate renal function.

All of the following criteria must be met:
- The patient did not progress during treatment with bortezomib
- If previously treated with lenalidomide, the patient did not discontinue lenalidomide due to adverse events and/or did not experience disease progression during the first 3 months of treatment
- If the patient was most recently treated with lenalidomide, the patient disease has not progressed at any time during treatment.

**Duration of Approval:** 1 Year

4. **Lenalidomide within a triplet combination with Daratumumab + Dexamethasone**

**Eligibility Criteria:**
- Lenalidomide in combination with daratumumab and dexamethasone for patients with relapsed multiple myeloma who meet the funding criteria for daratumumab through the NDFP.
- Treatment should be in patients who have good performance status who have received at least one prior therapy.
- All cycles of daratumumab are given in combination with lenalidomide and dexamethasone as part of an every 4-week treatment cycle.
- Patients who were previously treated with lenalidomide or are currently on lenalidomide will only be eligible for the addition of daratumumab if lenalidomide was not previously discontinued due to adverse events and the patient’s disease is not refractory\(^1\) to lenalidomide.

\(^1\)Refractory disease is defined as:
- Disease progression within 60 days of any dose of lenalidomide, or
- Disease progression while on lenalidomide, or
- Failure to achieve at least a minimal response while on lenalidomide.
Lenalidomide
Brand(s): Revlimid and generics (see formulary for funded OFIs)
DOSAGE FORM/ STRENGTH: 2.5mg, 5 mg, 10 mg, 15 mg, 20 mg, 25 mg capsule

Treatment with lenalidomide within a triplet combination with daratumumab should be continued until disease progression or unacceptable toxicity to daratumumab or to lenalidomide.

Renewals will be considered for patients who continue to respond to a triple therapy regimen with daratumumab and dexamethasone. (i.e. Is not relapsed/refractory)

Duration of Approval of Initials and Renewals: 1 year

Lenvatinib
Brand(s): Lenvima
DOSAGE FORM/ STRENGTH: 4 mg, 10 mg capsules (packaged as 8mg, 10mg, 14mg, 18 mg, 20 mg, and 24 mg daily dose cartons.
Updated March 5, 2020

For the treatment of patients with locally recurrent or metastatic, progressive, differentiated thyroid cancer (DTC) who meet ALL the following criteria:

- Papillary or Follicular subtypes of DTC that are histologically or cytologically confirmed; AND
- Thyroid cancer is refractory or resistant to radioactive iodine; AND
- DTC shows evidence of disease progression within the past 13 months; AND
- Patient has good performance status with ECOG less than or equal to 2; AND
- Lenvatinib is being used as monotherapy

Exclusion criteria:
- Patients with anaplastic or medullary thyroid cancer
- Patients who have received more than one prior therapy with a tyrosine kinase inhibitor

Duration of Approval: 1 Year

Renewal of funding will be considered until a patient progresses on treatment or develops unacceptable toxicity to lenvatinib.

Duration of Approval: 1 Year
Lenvatinib
Brand(s): Lenvima
DOSAGE FORM/ STRENGTH: 4 mg, 10 mg capsules (packaged as 8mg, 10mg, 14mg, 18 mg, 20 mg, and 24 mg daily dose cartons.

For the treatment of unresectable advanced\(^1\) hepatocellular carcinoma (HCC) in adult patients who meet ALL the following criteria prior to starting treatment with lenvatinib;

- Patient is 18 years of age or older; AND
- Lenvatinib will be used as monotherapy for HCC; AND
- Patient has good performance status with Eastern Cooperative Oncology Group (ECOG) Performance status less than or equal to 2; AND
- Has a Child-Pugh class A liver function.

\(^1\) Patients with Stage B HCC, based on the Barcelona Clinic Liver Cancer (BCLC) Staging System will be considered for lenvatinib if they have progressed on transarterial chemoembolization (TACE). Case-by-case consideration will be provided for Stage B HCC patients who are not suitable for the TACE procedure. In such situations, please provide additional information to support why the patient is not suitable for TACE.

Exclusion Criteria: Patients meeting any of the following criteria will not be funded.

- Patients with Child-Pugh score greater than 6 (i.e. Child-Pugh class B or C) will not be funded.
- Patients who have progressed on sorafenib for HCC will not be funded for lenvatinib

Only one of sorafenib or lenvatinib for the treatment of HCC will be funded in the first line. Patients will be permitted to switch from sorafenib to lenvatinib if they experience intolerance and have not progressed on sorafenib.

Recommended Dosage:

The recommended daily dose of lenvatinib is 8mg once daily for patients with a body weight of <60kg and 12 mg once daily for patients with a body weight of ≥60 kg

Renewal Criteria:

Renewals will be considered for patients who have not experienced unacceptable toxicities to lenvatinib or until disease progression.

Please provide radiographic results, scan results or both indicating no progression. Progression evaluation will be in accordance with modified Response Evaluation Criteria in Solid Tumors (mRECIST) or RECIST 1.1 criteria.

Approval duration for initials and approvals: 3 months
For the treatment of adult patients diagnosed with FMS-like tyrosine kinase 3 (FLT3)-mutated acute myeloid leukemia (AML) who meet ALL the following criteria:

- FLT3 mutation is confirmed by an approved test; AND
- Midostaurin is used as first-line\(^1\) for FLT3-mutated AML; AND
- Midostaurin is used in combination with standard induction chemotherapy with cytarabine and daunorubicin followed by standard consolidation chemotherapy with cytarabine OR any 7+3 induction regimen containing idarubicin followed by standard consolidation chemotherapy with cytarabine.

Exclusion criteria:

Midostaurin will not be funded in the following situations:

- As maintenance therapy for AML;
- Patients who have developed therapy-related AML after radiation therapy or chemotherapy for another cancer or disorder;
- Patients receiving other induction chemotherapy regimens aside from those mentioned in the eligibility criteria or upon finishing the consolidation phase of treatment;
- Patients undergoing re-induction and/or re-consolidation.

Recommended Dose(s):

Induction dose: Midostaurin 50 mg twice daily on Days 8 to 21 with each cycle of induction cytarabine and daunorubicin.

A maximum of 2 induction cycles may be funded. (Note: EAP only considers funding of outpatient midostaurin usage.)*

Consolidation phase: Midostaurin 50 mg twice daily on days 8 to 21 of each cycle of consolidation with cytarabine.

A maximum of 4 consolidation cycles may be funded by EAP (for cycles administered as an outpatient)*.

Up to 2 cycles of induction and 4 cycles of consolidation may be funded in accordance with patient response to therapy.*
**Midostaurin**  
**Brand(s):** Rydapt  
**DOSAGE FORM/ STRENGTH:** 25 mg capsule

**Approval duration:** Up to 6 months (maximum of 2 cycles of induction and 4 cycles of consolidation) *

1 For a short-term, time limited period, the Ministry will consider requests from prescribers who wish to add midostaurin to their patients’ current regimens that have been initiated prior to the provincial funding of midostaurin. To be considered, patients must currently be on standard induction and consolidation chemotherapy, and have not experienced disease progression or unacceptable intolerance during the first-line treatment with the standard chemotherapies being used.

*EAP will provide coverage for midostaurin administered in the outpatient setting (e.g., consolidation cycles). Those responding to induction therapy may require ongoing access to midostaurin for the consolidation phase of treatment upon discharge from the hospital.

**Nilotinib**  
**Brand(s):** Tasigna  
**DOSAGE FORM/ STRENGTH:** 150 mg, 200 mg capsule

**For the treatment of patients with chronic phase Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML).**

*Note: Ministry will only fund any TWO of the oral Tyrosine Kinase inhibitors (TKIs) * used for chronic phase CML per patient in a lifetime. (* TKIs: imatinib, nilotinib, or dasatinib)*

*If the patient develops grade 3 or 4 toxicity on one of the above TKI’s within 3 months of initiating therapy, access to a 3rd oral TKI will be funded for that patient.*

Approved dose: 300 mg twice daily but not exceeding 800 mg/day

**Duration of Approval:** 1 Year
Nilotinib
Brand(s): Tasigna
DOSAGE FORM/ STRENGTH: 150 mg, 200 mg capsule

For the treatment of patients with accelerated phase Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML) with documented intolerance¹ or resistance² to imatinib therapy.

¹Intolerance to imatinib at any dose occurs where the Patient has experienced persistent grade 3 or grade 4 toxicity requiring discontinuation of imatinib therapy; or

²Imatinib resistance occurs where the Patient has primary or acquired resistance to imatinib at doses of at least 600mg/day or via a mutational analysis report.

Exclusion Criteria – Patients with the following exclusion criteria will not be funded:

- blast phase CML;
  - a. for Ph+ acute lymphocytic leukemia (ALL);
  - b. combination treatment with any two or more oral TKIs’s (imatinib, nilotinib, or dasatinib) will not be funded
  - c. For accelerated phase CML, nilotinib is not funded as a sequential third line therapy in patients who experience primary or acquired resistance (not including mutational resistance) to dasatinib.

Approved dosage: Up to 800 mg/day but doses above 800 mg per day will not be considered

Renewal Criteria:

Renewals are considered for patients who experience hematologic and/or cytogenetic response to therapy, is expected to continue to do benefit from therapy with Tasigna.

Duration of Approval: 1 Year
Initiation Criteria:

For the maintenance treatment of newly diagnosed or recurrent high grade epithelial ovarian, fallopian tube, or primary peritoneal cancer in adult patients who meet the following criteria;

1. Patients are diagnosed with high grade serous or endometrioid tumours classified as stage III or IV disease according to the International Federation of Gynecology and Obstetrics (FIGO) criteria at the time of their initial diagnosis; AND

2. Patient is using niraparib as maintenance therapy immediately after one course of first line platinum-based therapy in which radiological response (complete or partial) is demonstrated after completing between 6 and 9 cycles of treatment,\(^1\) OR

Patient is using niraparib as maintenance therapy in recurrent disease after having received at least two prior courses of chemotherapy in which platinum sensitive disease\(^2\) was demonstrated with at least one completed prior treatment course AND radiological response (complete or partial) must be demonstrated after completing at least 4 cycles of their most recent course of chemotherapy\(^3\);

AND

3. Patients using niraparib as maintenance as first line or in the recurrent/relapsed setting must initiate niraparib within 12 weeks of the final dose of the chemotherapy;\(^4\) AND

4. Niraparib must be used as monotherapy for maintenance therapy; AND

5. Patient has good performance status.

\(^1\)Case-by-case consideration will be provided for newly diagnosed patients who are unable to complete 6 cycles of a platinum-based treatment OR who are unable to use a platinum-based chemotherapy but have achieved complete or partial radiological response after treatment with alternative, non-platinum-based chemotherapy.

\(^2\)Platinum-sensitive disease is defined as disease progression/recurrence/relapse occurring at least 6 months following completion of a platinum-based chemotherapy in which an initial response had been demonstrated.
Niraparib
Brand(s): Zejula
DOSAGE FORM/ STRENGTH: 100 mg capsule

Case-by-case consideration will be provided for patients with recurrent disease who are unable to use a platinum-based chemotherapy after having demonstrated platinum-sensitive disease to an earlier line of treatment after achieving complete or partial radiological response with their most recent course of treatment with alternative, non-platinum-based chemotherapy.

If more than 8 weeks have elapsed from the last chemotherapy treatment, consideration should be given to exclude disease progression before start of therapy.

Exclusion Criteria:

1. Patients who have previously progressed while on niraparib maintenance therapy will not be funded.
2. Patients who have experienced disease progression while on treatment with a prior PARP inhibitor regardless of treatment line (i.e. first line or recurrent/relapsed setting) will not be funded for niraparib.
3. Patients who have developed disease progression before start of niraparib maintenance therapy will not be funded.
4. Patients with symptomatic, uncontrolled brain metastases will not be funded.
5. Niraparib is not funded when used as combination therapy with chemotherapy.
6. Niraparib retreatment will not be funded in those who have completed 3 years of maintenance therapy and experienced disease progression.

Renewal Criteria:

Niraparib maintenance therapy after first line platinum-based treatment:
Ongoing funding will be considered until disease progression or development of unacceptable toxicity or up to a maximum of 3 years if there is no evidence of disease recurrence.

Niraparib maintenance therapy in recurrent platinum-sensitive disease:
Ongoing funding will be considered until disease progression or development of unacceptable toxicity

Notes:
1. Niraparib funding will not be considered for patients with histologies other than serous or endometroid unless the patient is BRCA mutated. Patients presenting with non-mucinous histology should be BRCA tested.
2. Imaging to rule out disease progression is required for patients delayed in starting maintenance therapy with niraparib. (Note: CA-125 clinical assessments may be considered case-by-case where imaging is not available).

3. Cancer antigen 125 (CA-125) and clinical assessments should be done at least every 3 to 4 months to monitor for disease reoccurrence or progression.

4. Patients with no evidence of disease after chemotherapy who are initiated niraparib in the first line maintenance setting will be funded for a maximum of 3 years.

5. Time-limited access to niraparib will be provided for patients already on maintenance bevacizumab who wish to switch to niraparib monotherapy as long as other criteria are met and there is no evidence of disease progression.

6. Patients on another PARP treatment may switch to niraparib if they have developed intolerances or allergies and if they have not experienced disease progression.

7. Niraparib will not be funded as maintenance therapy in patients who have previously progressed on niraparib or another PARP used as maintenance therapy after one or more lines of chemotherapy.

8. Retreatment with maintenance niraparib will not be funded in patients who previously completed 3 years of niraparib maintenance therapy after first line platinum-based chemotherapy, and then subsequently experienced disease progression.

9. Patients who have used a non-PARP maintenance therapy after first line chemotherapy may be funded for niraparib as maintenance in a subsequent line as long as disease progression had not occurred for 6 months after stopping maintenance therapy with the prior treatment AND upon meeting the initiation criteria again. Note that the ministry will only fund one PARP therapy (i.e. niraparib or olaparib) in the maintenance setting upon meeting all initiation criteria.

**Recommended dose:** 200 mg to 300 mg daily (Refer to the product monograph for dosing information.)

**Approval duration of initials and renewals:** 1 year

Reimbursement will be provided up to a maximum of 3 years if there is no evidence of disease recurrence.
Olaparib
Brand(s): Lynparza
DOSAGE FORM/ STRENGTH: 50mg capsules (Restricted access for patients already using capsules), 100mg tablets, 150mg tablets
Updated December 23, 2020

For the maintenance treatment of BRCA-mutated, high grade epithelial ovarian, fallopian tube, or primary peritoneal cancer in adult patients who meet ALL the following criteria;

1. Patient has documented mutation in BRCA1 or BRCA2 genes (germline or somatic detected by an approved testing method); AND

2. Patient is using olaparib as maintenance therapy immediately after one course of first line platinum-based chemotherapy in which radiological response (complete or partial) is demonstrated after at least 4 cycles of treatment; OR

   Patient is using olaparib as maintenance therapy in relapsed disease after having received more than one prior course of platinum-based chemotherapy in which platinum sensitive disease\(^1\) was demonstrated with one completed treatment course, and there is radiologic response (complete or partial) to the most recently completed course of platinum-based treatment.\(^2\)

   AND

3. Olaparib is started within 8 weeks of the patient’s final dose of platinum-based chemotherapy\(^2\) or within 12 weeks\(^3\) with restaging to confirm no disease progression if delay of more than 8 weeks since last dose of chemotherapy has occurred; AND

4. Olaparib is being used as monotherapy for maintenance treatment; AND

5. Patient has good performance status.

\(^1\)Platinum-sensitive disease is defined as disease progression/recurrence/relapse occurring at least 6 months following completion of a platinum-based chemotherapy in which an initial response had been demonstrated.

\(^2\)Patients who are unable to use a platinum-based chemotherapy after having demonstrated platinum- sensitive disease to an earlier line of treatment may be considered on a case-by-case basis if they have received at least 4 cycles of a non-platinum treatment, submit documentation for clinically relevant allergies or intolerance to platinum treatment, and meet all other aspects of the above criteria.
Olaparib
Brand(s): Lynparza
DOSAGE FORM/ STRENGTH: 50mg capsules (Restricted access for patients already using capsules), 100mg tablets, 150mg tablets

3 Patients not able to start olaparib within 12 weeks due to extenuating circumstances may be considered on a case-by-case basis if they have no evidence of disease progression, provide information to explain why treatment could not be started within 12 weeks, and meet all other aspects of the above criteria.

Exclusion Criteria: (Patients meeting any of the below criteria will not be funded.)

- Patients who have relapsed after at least one course of platinum-based chemotherapy and have not demonstrated platinum-sensitive disease¹.
- Patients who have developed disease progression before start of olaparib maintenance therapy.
- Retreatment with olaparib as maintenance therapy.

Olaparib is not funded when used as combination with chemotherapy.

Recommended dose: 300 mg twice daily for oral tablets

Approval duration: 1 year

Notes:
1. Imaging to rule out disease progression is required for patients delayed in starting maintenance therapy with olaparib by more than 8 weeks or who have stopped therapy for more than 14 days prior to starting or restarting olaparib (Note: CA-125 clinical assessments may be considered case-by-case where imaging is not available).

2. Cancer antigen 125 (CA-125) and clinical assessments should be done at least every 3 to 4 months to monitor for disease reoccurrence or progression.

3. Olaparib will be funded for a maximum of 2 years in the maintenance setting after first line platinum-based therapy if there is no evidence of disease. Olaparib maintenance therapy will be funded ongoing until disease progression or development of unacceptable toxicity to olaparib for those using in relapsed platinum-sensitive disease.

4. Time limited access to olaparib will be provided for patients already on bevacizumab maintenance who wish to switch to olaparib monotherapy as long as other criteria are met and there is no evidence of disease progression on imaging and within 12 weeks of completing chemotherapy.
Olaparib
Brand(s): Lynparza
DOSAGE FORM/ STRENGTH: 50mg capsules (Restricted access for patients already using capsules), 100mg tablets, 150mg tablets

Renewal Criteria:

Olaparib maintenance therapy after first line platinum-based treatment.

Ongoing funding will be considered until disease progression or development of unacceptable toxicity or up to a maximum of 2 years if there is no evidence of disease.

Ongoing funding will be considered until disease progression or development of unacceptable toxicity

Recommended Dose: 300 mg twice daily for oral tablets

Approval duration of renewals: 1 year

(Note that Olaparib will be funded for a maximum of 2 years in the maintenance setting after first line platinum-based therapy if there is no evidence of disease.)
Osimertinib
Brand(s): Tagrisso
DOSAGE FORM/ STRENGTH: 80mg tablets (40mg on case-by-case)
Updated January 10, 2020

Initiation criteria:

For the treatment of locally advanced (not amenable to curative therapies) or metastatic non-small cell lung cancer (NSCLC) in individuals meeting the following criteria:

1. Previously untreated\(^1\) NSCLC in a patient with tumours that are documented to have Epidermal Growth Factor Receptor (EGFR) exon 19 deletions (exon 19 del) or exon 21 (L858R) substitution mutations (either alone or in combination with other EGFR mutations)

OR

Previously treated NSCLC in a patient who has experienced disease progression on one EGFR tyrosine kinase inhibitor (TKI) therapy (i.e. afatinib, gefitinib or erlotinib) with tumours that are documented to have EGFR T790M resistance mutations\(^2\); AND

2. Has a good performance status; AND

3. Osimertinib is being used as monotherapy.

Exclusion Criteria:

- Patients with EGFR wild-type mutations
- Patients with EGFR unknown mutations
- Osimertinib will not be funded as a third-line TKI
- Patients with EGFR exon 19 deletions (exon 19 del) or exon 21 (L858R) substitution mutations who receive afatinib or gefitinib in first line are not eligible for osimertinib in the 2\(^{nd}\) line NSCLC setting.\(^3,4\)

Note:

\(^1\) Eligible patients should be previously untreated in the locally advanced or metastatic setting.

\(^2\) Patients with de novo EGFR 790M mutations may be considered case-by-case.

\(^3\) Time-limited consideration will be provided for patients meeting all the above criteria who are currently on a first-, or second-generation EGFR TKI (i.e. gefitinib, afatinib, erlotinib) who have not experienced disease progression or patients who are currently on chemotherapy and are found to harbour a sensitizing or resistance mutation who wish to switch to osimertinib therapy.
Osimertinib
Brand(s): Tagrisso
DOSAGE FORM/ STRENGTH: 80mg tablets (40mg on case-by-case)

4Patients who progress on osimertinib in the first line will not be considered for another targeted TKI therapy (i.e. gefitinib or afatinib) for NSCLC

Renewal of funding of osimertinib will be considered in patients who continue to derive benefit from treatment. (i.e. until clinically meaningful progression occurs or development of unacceptable toxicities.)

Recommended dose: 80 mg per day orally

(Note that 40mg tablets are approved on a case-by-case basis to ensure cost-effectiveness of the funded strength.)

Duration of Approval for Initial and Renewal requests: 6 months
Palbociclib
Brand(s): Ibrance
DOSAGE FORM/ STRENGTH: 75 mg, 100 mg, 125 mg tablet or capsules
Updated December 4, 2020

**Initial Criteria**

For the treatment of patients with estrogen receptor(ER)-positive, human epidermal growth factor receptor 2 (HER 2)-negative; unresectable locally advanced breast cancer or metastatic breast cancer in patients who meet the following criteria:

1. Palbociclib is being used as combination therapy in one of the following treatment regimens¹:
   i) As first line therapy in combination with an aromatase inhibitor (i.e. letrozole, anastrozole, or exemestane) or fulvestrant in a patient who has not progressed on a prior systemic treatment (i.e. chemotherapy, immunotherapy, or endocrine therapy) for their unresectable locally advanced or metastatic disease; OR
   
   ii) As second line therapy in combination with an aromatase inhibitor (i.e. letrozole, anastrozole, or exemestane) or fulvestrant after progression on a chemotherapy for unresectable locally advanced or metastatic disease; OR
   
   iii) As a second or subsequent line therapy in combination with fulvestrant after progression on any number of endocrine monotherapies with the exception of progression during prior fulvestrant therapy.

¹Note: EAP funding will be considered for only one CDK 4/6 inhibitor regimen (i.e. Palbociclib or Ribociclib) OR Everolimus based regimen for the treatment of unresectable locally advanced or metastatic disease. No funding for sequential treatment regimens involving palbociclib or ribociclib or everolimus will be considered.

AND

2. Patients who received anastrozole or letrozole in the neo-adjuvant or adjuvant setting, must demonstrate a minimum disease free interval of twelve (12) months after stopping therapy to qualify for funding of palbociclib in combination with anastrozole or letrozole. (Note: This does not apply to patients receiving tamoxifen or exemestane in the neoadjuvant or adjuvant setting who progress or relapse early on those treatments.)

3. Patient has good performance status defined as an Eastern Cooperative Oncology Group (ECOG) score of 0 to 2; AND

4. Patient does not have active or uncontrolled metastases to the central nervous system; AND
5. In the case of a Patient who is pre-menopausal or peri-menopausal, the Patient is receiving a luteinizing hormone-releasing hormone (LHRH) agonist to achieve chemically-induced menopause (Note: Women who have had an oophorectomy are considered to be post-menopausal); AND

6. The Patient has not experienced disease progression on any of the following regimens for locally advanced or metastatic breast cancer:
   (i) a palbociclib or ribociclib regimen;
   (ii) an everolimus regimen; or
   (iii) another CDK 4/6 regimen that has been publicly funded.

Renewal Criteria:

Renewals will be considered in patients who have not demonstrated evidence of disease progression or development of unacceptable toxicity requiring discontinuation while on palbociclib.

Exclusion Criteria (Note that exclusion criteria apply to both Initial eligibility criteria and renewal criteria.) Patients meeting the following criteria will not be funded.

i) Patient is using palbociclib as retreatment after disease progression on a prior palbociclib-based regimen in advanced breast cancer.

ii) Patient is using palbociclib with other drugs or in combinations other than those situations mentioned in 1 i), ii), iii) of the eligibility criteria.

iii) Patient is using palbociclib in combination with letrozole or anastrozole in the metastatic setting but has experienced progression in the neoadjuvant or adjuvant setting occurring during treatment or within 12 months of stopping treatment with letrozole or anastrozole;

iv) Patient is pre- or peri-menopausal who is not being treated with a luteinizing hormone-releasing hormone (LHRH) agonist.

v) Patient who is intending to use palbociclib with fulvestrant who has progressed on prior fulvestrant used as monotherapy or as part of another regimen.

vi) Patient whose disease has progressed during treatment with a ribociclib regimen, an everolimus regimen, or another CDK 4/6 inhibitor regimen used for advanced, metastatic breast cancer, unless that use was through a clinical trial.

vii) Patient who has active or uncontrolled central nervous system (CNS) metastases.

viii) The Patient is requesting Ibrance for use with fulvestrant and has extensive, symptomatic, potentially life-threatening visceral metastases.
On a time-limited basis, funding will be considered for the following on a case-by-case basis:

1. Patients who missed the opportunity to use palbociclib in the advanced setting in patients started on first-line, monotherapy with an aromatase inhibitor (AI) (e.g. letrozole, anastrozole, exemestane) AND who have not have not experienced disease progression with current AI therapy AND who meet the disease-free time requirement if anastrozole or letrozole was used previously in the adjuvant or neoadjuvant setting and the EAP request is submitted between the dates of December 4, 2020 to March 4, 2021.

2. Addition of palbociclib for patients already on fulvestrant in first, second or subsequent line who has not experienced disease progression on fulvestrant and who are CDK 4/6 inhibitor naïve and otherwise eligible for this therapy if the EAP request is submitted between the dates of December 4, 2020 to March 4, 2021.

3. A switch to palbociclib + fulvestrant at progression for patients already on endocrine/hormonal therapy other than fulvestrant and who are CDK4/6 naïve and otherwise meet the eligibility requirements for this therapy.

4. A switch to palbociclib + fulvestrant at progression for patients already on and benefitting from everolimus + exemestane, provided that the start of everolimus + exemestane was prior to December 4, 2020. Patients must be CDK 4/6 inhibitor naïve and otherwise eligible for this therapy.

**Dosing:**

Palbociclib (Ibrance) 125mg orally once a day for 21 consecutive days, followed by 7 days off treatment, in a combination regimen with one of the following:

- A continuous daily aromatase inhibitor or
- Fulvestrant 500mg administered intramuscularly on days 1 and 15 of cycle 1 and then on day 1 of each subsequent 28-day cycle.
Pazopanib
Brand(s): Votrient
DOSAGE FORM/ STRENGTH: 200 mg tablet

For first-line treatment of advanced or metastatic renal cell carcinoma of clear cell histology in patients with good performance status (ECOG* ≤ 1)

ECOG = Eastern Cooperative Oncology Group Performance Status

The approved dosage is 800 mg once daily.

Duration of Approval: 1 year

Renewals will be considered for patients who have benefited from therapy (i.e. no disease progression) and are expected to continue to do so. Exclusion criteria: Funding for Votrient will not be approved for patients who demonstrate disease progression while on sunitinib, sorafenib, temsirolimus, everolimus or other drugs approved for treatment of metastatic renal cell carcinoma.

Duration of Approval: 1 year
Pomalidomide
Brand(s): Pomalyst
DOSAGE FORM/ STRENGTH: 1 mg, 2 mg, 3 mg, 4 mg capsules

For the treatment of patients with relapsed and/or refractory multiple myeloma (MM) who meet the following criteria;

i) Patient has failed lenalidomide\(^1\); AND
ii) Patient has previously failed\(^2\) OR may have a contraindication OR demonstrated an intolerance\(^3\) to bortezomib; AND
iii) Patient has demonstrated disease progression following the last treatment\(^4\) used for MM.

\(^1\)Failure to lenalidomide may include failure to treatment received during the maintenance setting.

\(^2\)Failure to bortezomib can include patients who have received a course of bortezomib during which there was no disease progression, however, at the time of relapse of the patient’s MM, the patient is no longer eligible for retreatment with bortezomib.

\(^3\)Details of the patient’s intolerance(s) to bortezomib must be provided on the funding application to EAP.

\(^4\)The patient’s last treatment may be a regimen other than one containing lenalidomide or bortezomib.

**Duration of Approval: 1 year**

**Renewals** will be considered for patients who continue to respond to therapy (i.e. is not refractory and has not relapsed).

**Duration of Approval: 1 year**
Ponatinib  
Brand(s): Iclusig  
DOSAGE FORM/ STRENGTH: 15 mg, 45 mg tablets

**Chronic Phase CML:**

a. For the treatment of patients with Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML) in chronic phase and documented T315i mutation; OR

b. For the treatment of patients with Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML) in chronic phase with documented resistance/disease progression or intolerance to at least 2 prior oral TKIs (imatinib, dasatinib or nilotinib), where ponatinib would be the third or fourth line TKI.

**Duration of Approval:** 1 year

**Accelerated Phase CML:**

- For the treatment of patients with Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML) in accelerated phase and documented T315i mutation; OR

- For the treatment of patients with Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML) in accelerated phase with documented resistance/disease progression or intolerance to at least 2 prior oral TKIs (imatinib, dasatinib or nilotinib), where ponatinib would be the third or fourth line TKI.

**Duration of Approval:** 1 year

**Blast Phase CML:**

(a) For the treatment of patients with Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML) in blast phase and documented T315i mutation; OR

(b) For the treatment of patients with Philadelphia chromosome positive (Ph+) chronic myelogenous leukemia (CML) in blast phase with documented resistance/disease progression or intolerance to at least 2 prior oral TKIs (imatinib and dasatinib), where ponatinib would be the third or fourth line TKI.
**Ponatinib**
Brand(s): Iclusig
DOSAGE FORM/ STRENGTH: 15 mg, 45 mg tablets

For Philadelphia chromosome positive acute lymphoblastic leukemia (Ph+ALL):

a. For the treatment of patients with Philadelphia chromosome positive acute lymphoblastic leukemia (Ph +ALL) and documented T315i mutation; OR

b. For the treatment of patients with Philadelphia chromosome positive acute lymphoblastic leukemia (Ph +ALL) with documented resistance/disease progression or intolerance to imatinib and dasatinib, where ponatinib would be the third line TKI.

Renewals will be considered upon confirmation from the clinician that the patient has experienced hematologic and/or cytogenic response and is expected to continue to do so.

Duration of Approval: 1 year

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**Regorafenib**
Brand(s): Stivarga
DOSAGE FORM/ STRENGTH: 40 mg tablets
Updated November 18, 2020

For the treatment of metastatic and/or unresectable gastrointestinal stromal tumors (GIST) in patients who have had disease progression on, or intolerance to, imatinib and sunitinib

Dosage: 160 mg once daily for 3 weeks followed by 1 week of no therapy to comprise a cycle of 4 weeks.

Duration of Approval: 6 Months

Reimbursement of Stivarga will be considered as long as benefit is observed or until unacceptable toxicity occurs.

Renewals will be considered in patients who continue to derive benefit from therapy.

Duration of Approval: 6 Months
Regorafenib
Brand(s): Stivarga
DOSAGE FORM/ STRENGTH: 40 mg tablet

Initial Criteria:

For the treatment of unresectable, advanced hepatocellular carcinoma (HCC) in patients who meet ALL the following criteria prior to starting treatment with regorafenib;

- Patient is 18 years of age or older; AND
- Regorafenib will be used as monotherapy for HCC; AND
- Regorafenib will be used as second line therapy in a patient who has experienced disease progression during treatment with sorafenib or lenvatinib for HCC; AND
- If prior treatment was sorafenib, patient must have tolerated at least 400 mg a day for 20 days or more in the 28 days before stopping treatment with sorafenib; AND
- Has an Eastern Cooperative Oncology Group (ECOG) Performance status less than or equal to 1; AND
- Has a Child-Pugh class A liver function.

Exclusion Criteria: Patients meeting any of the following criteria will not be funded.

- Regorafenib will not be funded in the first line setting.
- Patients with Child-Pugh score greater than 6 (i.e. Child-Pugh class B or C) will not be funded.

Recommended Dosage: 160 mg once daily for 3 weeks followed by 1 week of no therapy to comprise a cycle of 4 weeks.

Renewal Criteria:

Renewals will be considered until disease progression (1) or until patient develops unacceptable toxicity. Please provide radiographic and/or scan results indicating no progression\(^2\) with requests for renewal of funding.

\(^1\) Evaluation according to modified Response Evaluation Criteria in Solid Tumors (mRECIST) or RECIST 1.1 criteria.

\(^2\) Approval duration for initials and renewals: 3 months
Initial Criteria:

For the treatment of patients with estrogen receptor(ER)-positive, human epidermal growth factor receptor 2 (HER 2)-negative; unresectable locally advanced breast cancer or metastatic breast cancer in patients who meet the following criteria;

1. Ribociclib is being used as combination therapy in one of the following treatment regimens¹:
   
   i) As first line therapy in combination with an aromatase inhibitor (i.e. letrozole, anastrozole, or exemestane) or fulvestrant in a patient who has not progressed on a prior systemic treatment (i.e. chemotherapy, immunotherapy, or endocrine therapy) for their unresectable locally advanced or metastatic disease; OR
   
   ii) As second line therapy in combination with an aromatase inhibitor (i.e. letrozole, anastrozole, or exemestane) or fulvestrant after progression on a chemotherapy for unresectable locally advanced or metastatic disease; OR

   iii) As a second or subsequent line therapy in combination with fulvestrant after progression on any number of endocrine monotherapies with the exception of progression during prior fulvestrant therapy.

¹Note: EAP funding will be considered for only one CDK 4/6 inhibitor regimen (i.e. Ribociclib or Palbociclib) OR Everolimus based regimen for the treatment of unresectable locally advanced or metastatic disease. No funding for sequential treatment regimens involving ribociclib or palbociclib or everolimus will be considered.

AND

2. Patients who received anastrozole or letrozole in the neo-adjuvant or adjuvant setting, must demonstrate a minimum disease free interval of twelve (12) months after stopping therapy to qualify for funding of ribociclib in combination with anastrozole or letrozole. (Note: This does not apply to patients receiving tamoxifen or exemestane in the neoadjuvant or adjuvant setting who progress or relapse early on those treatments.)

3. Patient has good performance status defined as an Eastern Cooperative Oncology Group (ECOG) score of 0 to 2; AND

4. Patient does not have active or uncontrolled metastases to the central nervous system; AND
Ribociclib
Brand(s): Kisqali
DOSAGE FORM/ STRENGTH: 200 mg tablets

5. In the case of a Patient who is pre-menopausal or peri-menopausal, the Patient is receiving a luteinizing hormone-releasing hormone (LHRH) agonist to achieve chemically-induced menopause (Note: Women who have had an oophorectomy are considered to be post-menopausal); AND

6. The Patient has not experienced disease progression on any of the following regimens for locally advanced or metastatic breast cancer:
   i. a ribociclib or palbociclib regimen;
   ii. an everolimus regimen; or
   iii. another CDK 4/6 regimen that has been publicly funded.

Renewal Criteria:

Renewals will be considered in patients who have not demonstrated evidence of disease progression or development of unacceptable toxicity requiring discontinuation while on ribociclib

Exclusion Criteria (Note that exclusion criteria apply to both Initial eligibility criteria and renewal criteria.)

Patients meeting the following criteria will not be funded.

   i) Patient is using ribociclib as retreatment after disease progression on a prior ribociclib-based regimen in advanced breast cancer.
   ii) Patient is using ribociclib with other drugs or in combinations other than those situations mentioned in 1 i), ii), iii) of the eligibility criteria.
   iii) Patient is using ribociclib in combination with letrozole or anastrozole in the metastatic setting but has experienced progression in the neoadjuvant or adjuvant setting occurring during treatment or within 12 months of stopping treatment with letrozole or anastrozole;
   iv) Patient is pre- or peri-menopausal who is not being treated with a luteinizing hormone-releasing hormone (LHRH) agonist.
   v) Patient who is intending to use ribociclib with fulvestrant who has progressed on prior fulvestrant used as monotherapy or as part of another regimen.
   vi) Patient whose disease has progressed during treatment with a palbociclib regimen, an everolimus regimen, or another CDK 4/6 inhibitor regimen used for advanced, metastatic breast cancer, unless that use was through a clinical trial.
   vii) Patient who has active or uncontrolled central nervous system (CNS) metastases.
Ribociclib
Brand(s): Kisqali
DOSAGE FORM/ STRENGTH: 200 mg tablets

On a time-limited basis, funding will be considered for the following on a case-by-case basis:

1. Patients who missed the opportunity to use ribociclib in the advanced setting in patients started on first-line, monotherapy with an aromatase inhibitor (AI) (e.g. letrozole, anastrozole, exemestane) AND who have not have not experienced disease progression with current AI therapy AND who meet the disease-free time requirement if anastrozole or letrozole was used previously in the adjuvant or neoadjuvant setting and the EAP request is submitted between the dates of November 18, 2020 to February 18, 2021.

2. Addition of ribociclib for patients already on fulvestrant in first, second or subsequent line who has not experienced disease progression on fulvestrant and who are CDK 4/6 inhibitor naïve and otherwise eligible for this therapy if the EAP request is submitted between the dates of November 18, 2020 to February 18, 2021.

3. A switch to ribociclib + fulvestrant at progression for patients already on endocrine/hormonal therapy other than fulvestrant and who are CDK4/6 naïve and otherwise meet the eligibility requirements for this therapy.

4. A switch to ribociclib + fulvestrant at progression for patients already on and benefitting from everolimus + exemestane, provided that the start of everolimus + exemestane was prior to November 18, 2020. Patients must be CDK 4/6 inhibitor naïve and otherwise eligible for this therapy.

Dosing:
Ribociclib (Kisqali) 600mg orally once daily for 21 consecutive days, followed by 7 days off treatment in combination with one of the following:
- A continuous daily aromatase inhibitor or
- Fulvestrant 500mg administered intramuscularly on days 1 and 15 of cycle 1 and then on day 1 of each subsequent 28-day cycle.

Approval duration of Initials and Renewals: 1 year
Ruxolitinib  
Brand(s): Jakavi  
DOSAGE FORM/ STRENGTH: 5 mg, 10mg, 15 mg, 20 mg tablets

For the treatment of intermediate to high risk symptomatic Myelofibrosis (MF) in patients meeting the following criteria:

i) MF is assessed using the Dynamic International Prognostic Scoring System (DIPSS) Plus; or the patient has symptomatic splenomegaly

ii) Patient has an Eastern Cooperative Oncology Group (ECOG) performance status ≤ 3

iii) Patient is previously untreated or refractory to other treatment

Dosing regimen: 5 mg to 25 mg twice a day

Duration of Approval: 1 Year

Initial Renewals are considered for patients who:

- Have confirmation of either a reduction in spleen size or documented improvement of disease symptoms within 6 months of initiating therapy with Jakavi.

Second and subsequent Renewals are considered for patients who continue to benefit from therapy with Jakavi

For the treatment of patients with polycythemia vera who meet the following criteria:

a) Demonstrated resistance\(^1\) or demonstrated intolerance\(^2\) to hydroxyurea (HU); AND

b) Have a good performance status (ECOG ≤ 3)

\(^1\)Resistance to Hydroxyurea as defined by:

Use of HU for at least 3 months of treatment at a dose of at least 2 grams per day (or at maximally tolerated doses if unable to take 2 grams per day) meeting one of the following:

i. Patient continues to require phlebotomy to keep hematocrit (HCT) at less than 45%; OR

ii. Patient demonstrates uncontrolled myeloproliferation (i.e. platelet count > 400 x 10\(^9\)/L and white blood cell count > 10 x 10\(^9\)/L); OR

iii. Symptomatic splenomegaly\(^1\)Intolerance to Hydroxyurea as defined:

- After any dose of hydroxyurea, patient demonstrates one of the following:
  - Absolute neutrophil count < 1 x 10\(^9\)/L or platelet < 100 x 10\(^9\)/L or hemoglobin < 100 g/L at the lowest dose of HU required to achieve a response; OR
## Ruxolitinib

**Brand(s): Jakavi**  
**DOSAGE FORM/ STRENGTH:** 5 mg, 10 mg, 15 mg, 20 mg tablets

- Presence of leg ulcers or other unacceptable HU-related grade 3 or 4 non-hematological toxicities (e.g., mucocutaneous manifestations, gastrointestinal symptoms, pneumonitis, fever); OR
- If patient demonstrates non-hematological grade 2 toxicities for at least one week; OR
- If toxicity requires permanent discontinuation of HU, interruption of HU until resolution of toxicity, or requiring hospitalization as a result of HU toxicity.

### Renewal:

Patient continues to respond<sup>3</sup> to treatment and has not experienced disease progression.

<sup>3</sup>Response defined by any one or more of the following:
- Hematocrit <45% without phlebotomy
- Platelet count ≤ 400 x 10<sup>9</sup>/L
- White blood cell count ≤ 10 x 10<sup>9</sup>/L
- Non-palpable spleen

Approval duration for initiation and renewal: 1 year

## Sorafenib

**Brand(s): Nexavar**  
**DOSAGE FORM/ STRENGTH:** 200 mg tablet

For the treatment of metastatic renal cell carcinoma (MRCC) as second-line treatment for patients who have:

- a) Histologically confirmed metastatic clear-cell renal-cell carcinoma; AND
- b) Experienced disease progression after prior cytokine therapy within the previous 8 months; AND
- c) A performance status of 0 or 1 on the basis of the Eastern Cooperative Oncology Group criteria; AND
- d) Intermediate-risk or low-risk status, according to the Memorial Sloan-Kettering Cancer Center (MSKCC) prognostic score.

### Duration of Approval:

1 year

### Renewals

Will be considered with confirmation from the physician that the patient has benefited from therapy and is expected to continue to do so.
**Sorafenib**  
*Brand(s): Nexavar*  
*DOSAGE FORM/STRENGTH: 200 mg tablet*

For the treatment of **advanced hepatocellular carcinoma (HCC)** in patients who have:  
a) Child-Pugh Class A disease; and  
b) ECOG* status 0, 1 or 2; and  
c) Either progressed on transarterial chemoembolization (TACE) or are not suitable for the TACE procedure (where detailed rationale is provided).

**Duration of Approval:** 3 months

*ECOG = Eastern Cooperative Oncology Group Performance Status*

**Renewal** will be considered for patients with documentation of radiography and/or scan results indicating no diseases progression.

**Sunitinib**  
*Brand(s): Sutent*  
*DOSAGE FORM/STRENGTH: 12.5 mg, 25 mg, 50 mg capsule*

For the treatment of **gastrointestinal stromal tumour (GIST)** in patients with unresectable or metastatic/recurrent GIST where one of the following conditions is met:  
- Early progression (within 6 months) while on imatinib; OR  
- Progression following treatment with optimum (escalated) doses of imatinib (800mg per day); OR  
- Intolerance* to imatinib (where detailed description of intolerance is provided).

*Definition of intolerance to imatinib – patient has experienced persistent grade 3 toxicity requiring discontinuation of therapy.

**Duration of Approval:** 6 months

**Renewal** will be considered for patients who are stable (no disease progression) and not experiencing intolerance to sunitinib therapy.

*Note: Approval will be granted at a dose of 50mg per day (4 weeks on, 2 weeks off).*

For the treatment of **metastatic renal cell carcinoma (MRCC):**

- **First-line therapy** for patients with MSK Prognostic Score of Favourable Risk or an Intermediate Risk OR  
- **Second-line therapy** for patients where:  
  - The disease is of clear cell histology AND  
  - Documented failure to first-line cytokine-based therapy
Sunitinib
Brand(s): Sutent
DOSAGE FORM/ STRENGTH: 12.5 mg, 25 mg, 50 mg capsules

Duration of Approval: 1 year

Renewal will be considered for patients with documentation of radiography and/or scan results indicating no diseases progression.

Duration of Approval: 1 year

Note: The prescribed dosage should be 50 mg daily for four (4) weeks, followed by two (2) weeks off the Drug Product, in repeated six (6) week cycles.

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

Exclusion criteria: Sutent will not be approved for second-line sequential therapy after everolimus failure in the first-line setting.

Dosing: 37.5 mg daily

Thalidomide
Brand(s): Thalomid
DOSAGE FORM/ STRENGTH: 50 mg capsule, 100 mg capsule, 200 mg capsule

For the treatment of Multiple Myeloma in patients 65 years of age or older meeting the following criteria;

a) Thalidomide is being used in combination with melphalan and prednisone; AND
b) The patient has not previously received other treatments¹ for multiple myeloma; AND
c) The patient is deemed to be unsuitable for stem cell transplantation; AND

¹ Exception is for those meeting bortezomib criteria as described below.

It should be noted that funding of thalidomide will be considered on a case-by-case basis for patients who have developed severe (grade III/IV) thrombocytopenia during the first 1 to 2 cycles of treatment with bortezomib and who have not experienced disease progression on bortezomib.

Duration of Approval: A maximum of 12 six-week cycles

Exclusion criteria:

Funding will not be considered for patients who are using thalidomide as second-line treatment of multiple myeloma.
Initial criteria:

For the mutation-targeted treatment of patients with BRAF V600 mutation-positive unresectable melanoma or metastatic melanoma meeting the following criteria:

- As monotherapy or as combination therapy with dabrafenib.
- If brain metastases are present, they should be asymptomatic or stable

Exclusion Criteria:
- BRAF V600 negative, or wild type tumors, or unknown status will not be funded
- Funding will not be considered in patients who have experienced progression on a BRAF mutation targeted therapy. The Ministry will fund only one BRAF mutation targeted treatment/treatment regimen.
- May be sequenced after immunotherapies or other funded treatments, however, treatment beyond third line will not be considered for funding.

Recommended Dose as Monotherapy:

2 mg once daily until disease progression or development of unacceptable toxicity requiring discontinuation of trametinib

Recommended Dose as combination dual therapy with Dabrafenib:

Trametinib 2mg once daily and Dabrabenib 150 mg twice daily, until disease progression or development of unacceptable toxicity requiring discontinuation

Renewal criteria:

Therapy as monotherapy OR as combination dual therapy (as above) may be continued until evidence of disease progression¹ or development of unacceptable toxicity requiring discontinuation.

¹ Letter from physician outlining radiological and clinical benefit requiring continuation of the drug and verification of no disease progression must be submitted.

Approval duration (both initial and renewal requests): 6 months (patients should have their disease status assessed at least every 6 months)
**Trametinib**
**Brand(s):** Mekinist
**DOSAGE FORM/ STRENGTH:** 0.5 mg, 2 mg tablet

**Case by case:**

Requests in patients who have initiated another single-agent BRAF or MEK inhibitor therapy will be considered on a case-by-case basis ONLY IF there has been no disease progression.

Requests in patients who have initiated another single-agent BRAF or MEK inhibitor therapy will be considered on a case-by-case basis ONLY IF there has been no disease progression.

**Exclusion Criteria:**

- BRAF V600 negative, or wild type tumors, or unknown status will not be funded
- Trametinib therapy (as monotherapy or in combination with dabrafenib) will not be considered for funding in patients who have progressed on a prior BRAF inhibitor therapy used as monotherapy or in combination.

**For the adjuvant treatment of resected Stage III cutaneous melanoma in patients meeting ALL the following criteria;**

i) Trametinib is being used as combination therapy with Dabrafenib

ii) Patient's cutaneous melanoma met the following requirements prior to resection:
   a. Histologically confirmed stage IIIA (limited to lymph node metastases of > 1 mm), IIIB, IIIC, or IIID cutaneous melanoma
   [8th edition of the American Joint Committee on Cancer staging system]
   b. BRAF V600 mutated (all BRAF V600 mutations)

iii) Post-resection, clinical or radiographic confirmation of complete disease resection including absence of in-transit metastases must be provided.¹

iv) Eastern Cooperative Oncology Group (ECOG) Performance Status 0 to 2

¹Micrometastatic lymph node involvement detected by sentinel lymph node biopsy will be allowed.

**Exclusion Criteria:**

- Patients with stage IIIA cutaneous melanoma with lymph node metastases less than 1 mm.
- Monotherapy with Dabrafenib
- Combinations with other anticancer therapies
Trametinib
Brand(s): Mekinist
DOSAGE FORM/ STRENGTH: 0.5 mg, 2 mg tablet

Recommended Dose in adjuvant therapy:

Trametinib 2mg once daily and Dabrabenib 150 mg twice daily

Notes: Treatment administered post-resection until disease recurrence or unacceptable toxicity to a maximum of 12 months of treatment in total.

(Note: 12 months refers to duration of adjuvant treatment accessed through all sources of funding (i.e. private and public).

Approval duration: Maximum of 12 months. Renewals are not considered.

Additional notes:

1. The Ministry will reimburse for provincially funded treatments for use in the adjuvant setting in patients with cutaneous melanoma for a total duration of up to 12 months. Overall access to adjuvant therapy will be limited to 12 months in total and combines the duration of use of all treatments administered in the adjuvant setting.

2. Funding will not be granted for dabrafenif-trametinib in patients who have used another treatment in the adjuvant setting for cutaneous melanoma for a duration of 3 months or longer.

3. A one-time switch to dabrafenif-trametinib in the adjuvant cutaneous melanoma setting will be permitted for BRAF-mutated patients who switch from another adjuvant treatment that has been used for less than 3 months and upon meeting the above funding criteria.

4. Patients who experience disease progression within 6 months of completion of adjuvant BRAF therapy will not be funded for another BRAF targeted therapy.

5. Switching to cobimetinib-vemurafenib from dabrafenib-trametinib will not be permitted in BRAF-positive patients.
**Trifluridine/Tipiracil**  
**Brand(s):** Lonsurf  
**DOSAGE FORM/ STRENGTH:** 15 mg/6.14mg tablet, 20 mg/8.19mg tablet  
**Effective date:** March 25, 2021

**Initial Criteria:**  
In combination with best supportive care (BSC) for the treatment of metastatic gastric cancer (mGC) or adenocarcinoma of the gastroesophageal junction (GEJ) in adult patients who meet ALL of the following criteria;

1. Patient has histologically confirmed non-resectable, metastatic, gastric adenocarcinoma (including GEJ) as defined by the American Joint Committee of Cancer (AJCC) staging classification; AND
2. Patient has experienced disease progression\(^1\) and/or clinically relevant intolerances or allergies with at least two (2) prior lines of chemotherapy including a fluoropyrimidine, a platinum, and either a taxane or irinotecan-containing regimen;  
   OR  
   Patient has HER2 positive disease and has experienced disease progression\(^1\) and/or clinically relevant intolerances or allergies with at least two (2) prior lines of chemotherapy including a fluoropyrimidine, a platinum, and either a taxane or irinotecan-containing regimen, and at least one HER2/neu-targeted therapy;  
   AND
3. Patient has Eastern Cooperative Oncology Group (ECOG) status 0 or 1.  
\(^{1}\) As confirmed via imaging during/within 3 months of previously completed line(s) of therapy

Note: If patients have experienced intolerances with the prior lines of therapy, please provide a description of the intolerance including grade of toxicity which required the drug regimen to be stopped.

**Renewal Criteria:**  
Renewals will be considered until radiographic evidence of disease progression or development of unacceptable toxicities.

Renewals should include radiographic reports and will be evaluated according to modified Response Evaluation Criteria in Solid Tumors (mRECIST) or RECIST 1.1 criteria.

**Approval duration of initial and renewals:** 6 months.

**Recommended dose:**  
35 mg/m\(^2\)/dose twice daily on days 1 to 5 and days 8 to 12 of each 28-day cycle.  
Dose adjustments based on individual safety and tolerability.
Caprelsa (Vandetanib) is funded for the treatment of symptomatic and/or progressive¹ medullary thyroid cancer (MTC) in patients who meet the following criteria:

(a) Patient has unresectable locally advanced or metastatic disease; AND

(b) Vandetanib is being used as monotherapy for MTC; AND

(c) ECOG less than or equal to 2²; AND

(d) Prescribed by or in consultation with an oncologist or internist experienced with the treatment of MTC.

Exclusion Criteria:

(a) Patients with QT interval prolongation/abnormalities (e.g. QTc that is unmeasurable or greater than or the same as 480ms) or who are taking medications that prolong QT interval prolongation.

(b) Patients with indolent, asymptomatic, or slowly progressive disease.

(c) Vandetanib is not funded as combination therapy

Renewal Criteria:

Renewal of funding will be provided until disease progression or development of unacceptable toxicity³.

¹As confirmed by radiological reports.

²Patients with an ECOG greater than 2 will be considered case-by-case upon submission of information regarding the risk of toxicity.

³At the time of renewal, prescriber should address whether there have been any significant cardiac events or concerns regarding cardiovascular toxicities.

Duration of approval for initial and renewal criteria: 1 year

Note: Prescribers and dispensing pharmacies are presumed to be in compliance with the requirements of the Caprelsa Restricted Distribution Program which is administered through the manufacturer.
Initiation Criteria:

For the treatment of adult patients with relapsed refractory chronic lymphocytic leukemia (CLL) or small lymphocytic lymphoma (SLL) meeting the following criteria;

1. Venetoclax will be used as monotherapy meeting one of the following circumstances;
   i) As second line therapy in a patient with high risk cytogenic markers [chromosome 17p deletion, TP 53 mutation OR unmutated immunoglobulin heavy chain variable region (IgHV)] who has experienced disease progression to ibrutinib.

   ii) As third or fourth line therapy in a patient who has experienced disease progression to at least two prior treatment regimens in which one treatment must be a B-cell receptor inhibitor (BCRi), specifically ibrutinib OR idealisib in combination with rituximab.

   OR

2. Venetoclax will be used in combination with Rituximab administered intravenously\(^1\) meeting one of the following circumstances;
   i) As a second line therapy in a patient with high-risk cytogenic markers [chromosome 17p deletion, TP 53 mutation, OR unmutated immunoglobulin heavy chain variable region (IgHV)] who has experienced disease progression while on ibrutinib in the first line.

   ii) As second line therapy\(^2\) in a patient who has experienced disease progression with a chemotherapy ± immunotherapy\(^3\) (e.g. fludarabine based regimen ±rituximab, bendamustine, cyclophosphamide-vincristine-prednisone combination) regimen OR chlorambucil in combination with obinutuzumab in the first line.

   iii) As third line therapy\(^2\) in a patient who has experienced disease progression to two (2) prior treatments that may include one of the following:
      • Two chemotherapy ± immunotherapy\(^3\) regimens
      • A chemotherapy ± immunotherapy\(^3\) regimen in the first line and ibrutinib OR idealfilisib in combination with rituximab in the second line
      • Chlorambucil in combination with obinutuzumab in the first line followed by ibrutinib OR idealfilisib in combination with rituximab in the second line
Venetoclax
Brand(s): Venclexta
Dosage Form/Strength: 10 mg, 50 mg, 100 mg, Starter Pack (4 week supply)

DOSAGE FORM/ STRENGTH
iv) As fourth line therapy\(^2\) in a patient who has experienced disease progression to three (3) prior lines of treatment that may include any of the following circumstances:

- Two chemotherapy ± immunotherapy\(^3\) regimens followed by ibrutinib OR idelalisib in combination with rituximab in the third line
- Chlorambucil in combination with obinutuzumab in the first line followed by a chemotherapy ± immunotherapy\(^3\) regimen in the second line, and ibrutinib OR idelalisib in combination with rituximab in the third line.

3. Patient has good performance status with ECOG equal to or less than 2

Notes:
- Retreatment with venetoclax in combination with rituximab therapy may be considered at the time of relapse for patients able to tolerate and complete 24 months of a combination venetoclax and rituximab regimen and who have demonstrated a progression-free interval of at least 12 months after stopping therapy.
- Patients on venetoclax monotherapy who have been responding to treatment (i.e. have not progressed) but who are unable to achieve an adequate response may be considered for rituximab to be added to the treatment.
- Patients with severe intolerances to treatments required in prior lines of therapy will be considered case-by-case. Please provide details of the grade of toxicity experienced and/or clinical details to justify the contraindication to treatment options.

Renewals for venetoclax monotherapy will be considered until disease progression or development of unacceptable toxicity.

Renewals for venetoclax in combination with rituximab will be considered until disease progression or development of unacceptable toxicity up to a maximum of two years, whichever comes first.

Exclusion Criteria:
- Patient who has experienced disease progression while on venetoclax

- Venetoclax as monotherapy or as combination therapy with rituximab will not be funded in fifth line or beyond. (However, venetoclax in combination with rituximab may be considered for retreatment upon relapse of CLL after experiencing a 12 month treatment-free interval following treatment with a venetoclax and rituximab combination regimen in a prior line of therapy).
Venetoclax
Brand(s): Venclexta
Dosage Form/Strength: 10 mg, 50 mg, 100 mg, Starter Pack (4 week supply)

Exclusion Criteria (continued):

- Patients who are NOT CD20 antibody sensitive (i.e. an individual who has experienced a relapse of CLL within 12 months of stopping/completing a rituximab-containing regimen or an obinutuzumab-containing regimen for CLL) will not be reimbursed for a venetoclax combination regimen with rituximab.
- Patient who has had an allogenic stem cell transplant in the 12 months preceding the request for EAP coverage for venetoclax as monotherapy or as combination therapy with rituximab.
- Patients with active or uncontrolled autoimmune cytopenias.

Dosing Regimen of Venetoclax monotherapy:

Using the venetoclax (Venclexta) "starter pack" for the ramp up phase, dose at 20 mg once daily for 7 days, followed by 50 mg once daily for 7 days, followed by 100 mg once daily for 7 days, followed by 200 mg once daily for 7 days, followed by 400 mg once daily until disease progression or unacceptable toxicity.

Approved Duration for venetoclax monotherapy initiation and renewals: 1 year

Dosing Regimen for Venetoclax within a rituximab combination regimen:

Using the venetoclax (Venclexta) "starter pack" for the ramp up phase, dose at 20 mg once daily for 7 days (week 1), 50 mg once daily for 7 days (week 2), 100 mg once daily for 7 days (week 3), followed by 200 mg once daily for 7 days (week 4), followed by 400 mg once daily until disease progression or unacceptable toxicity up to a maximum of two years (24 months from Cycle 1 Day 2 of rituximab) whichever comes first.

Dosing Regimen for Rituximab within a venetoclax combination regimen:

375 mg/metre$^2$ Intravenously$^1$ on Day 1 of Cycle 1 (to be initiated after the patient has completed 5 weeks of ramp-up schedule with venetoclax and having received the 400 mg dose of venetoclax for 7 days), followed by 500 mg/metre$^2$ on Day 1 of Cycles 2 to 6, for a total of 6 infusions$^1$ of rituximab.

Approved Duration for venetoclax within a combination regimen with rituximab - Initiation: 1 year

Renewal duration for venetoclax within a combination regimen with rituximab: 1 year (total treatment duration funded is a maximum of 2 years from the date of addition of rituximab to the treatment regimen.)
Venetoclax
Brand(s): Venclexta
Dosage Form/Strength: 10 mg, 50 mg, 100 mg, Starter Pack (4 week supply)

1 The use of rituximab subcutaneous may only be considered in patients who are able to tolerate at least one full dose of intravenous infusion of rituximab during the first cycle. If a patient is unable to receive the full IV rituximab dose, continue subsequent cycles with rituximab IV until a full IV dose can be successfully given.

2 Venetoclax in combination with rituximab will not be considered if the patient has not been able to maintain a treatment free interval of at least 12 months after having used a rituximab-containing regimen or an obinutuzumab-containing regimen in the prior line of therapy. (i.e. Patients will only be deemed to be sensitive to a CD20 antibody if they have experienced a treatment-free interval of 12 months or longer following treatment with a rituximab-containing regimen and/or an obinutuzumab-containing regimen.)

3 Provincially funded chemotherapy and immunotherapy combination regimens for CLL can be considered based on the patient’s fitness/frailty and tolerability to targeted options. Refer to a list of funded chemoimmunotherapy regimens on the Cancer Care Ontario (CCO) website at https://www.cancercareontario.ca/en/search?nav-search=CLL&sort_by=search_api_relevance=&Apply

Vemurafenib
Brand(s): Zelboraf
DOSAGE FORM/STRENGTH: 240 mg tablet

Initiation requests:

For the treatment of patients with BRAF V600 mutation-positive unresectable stage III or stage IV melanoma.
- As monotherapy or as combination therapy with cobimetinib
- If brain metastases are present, they should be asymptomatic or stable

Exclusion Criteria:
- BRAF V600 negative, or wild type tumors, or unknown status will not be funded
- The Ministry will fund only one BRAF mutation targeted treatment/treatment regimen.
- May be sequenced after immunotherapies or other funded treatments, however, treatment beyond third line will not be considered for funding.
Vemurafenib
Brand(s): Zelboraf
DOSAGE FORM/ STRENGTH: 240 mg tablet

Renewal requests:

Therapy as monotherapy OR as combination dual therapy (as above) may be continued until evidence of disease progression\(^1\) or development of unacceptable toxicity requiring discontinuation.

\(^1\) Letter from physician outlining radiological and clinical benefit requiring continuation of the drug and verification of no disease progression or development of unacceptable toxicity must be submitted.

Approval duration (both initial and renewal requests): 6 months (patients should have their disease status assessed at least every 6 months)

Case by case:

Requests in patients who have initiated another BRAF and/or MEK inhibitor as monotherapy or combination therapy will be considered on a case-by-case basis ONLY IF there has been no disease progression.

**Recommended Dose as Monotherapy:**

960 mg twice daily until disease progression or development of unacceptable toxicity requiring discontinuation of vemurafenib.

**Recommended Dose as combination dual therapy with Cobimetinib:**

Cobimetinib 60mg once daily for 21 days, followed by seven days off treatment; AND Vemurafenib 960mg twice daily for 28 days. Both drugs are given until disease progression or unacceptable toxicity.
Vismodegib  
Brand(s): Erivedge  
DOSAGE FORM/ STRENGTH: 150 mg tablet

For the treatment of metastatic basal cell carcinoma (BCC) or locally advanced BCC (including patients with basal cell nevus syndrome, i.e. Gorlin syndrome) in patients who meet the following criteria;

- Patient must have measurable metastatic disease or locally advanced disease;  
AND  
- Patient's disease must be considered inoperable or inappropriate for surgery\(^1\);  
AND  
- Patient's disease must be considered inappropriate for radiotherapy\(^2\); AND  
- Patient is 18 years or age or older; AND  
- Patient has an ECOG ≤ 2

Dose: 150 mg orally once daily taken until disease progression or unacceptable toxicity. Requests must include the following information:

**Duration of Approval:** 1 Year

Physicians must provide rationale for why surgery AND radiation cannot be considered

- The request must include a surgical consult note that provides a preoperative/surgical evaluation why surgery is not appropriate for the patient;  
AND  
- A consult note as to why radiation therapy is not appropriate for the patient; AND  
- Both of the above evaluations must come from a physician who is not the requesting physician; AND  
- The request must include confirmation that the patient has been discussed at a multi-disciplinary cancer conference (MCC) or equivalent.

\(^1\)Considered inoperable or inappropriate for surgery for at least ONE of the following reasons:

- Technically not possible to perform surgery due to size/location/invasiveness of BCC (either lesion too large or can be several small lesions making surgery not feasible); OR  
- Recurrence of BCC after two or more surgical procedures and curative resection unlikely; OR  
- Substantial deformity and/or morbidity anticipated from surgery.

\(^2\)Considered inappropriate for radiation for at least ONE of the following reasons:

- Contraindication to radiation (e.g. Gorlin syndrome); OR  
- Prior radiation to lesion; OR  
- Suboptimal outcomes expected due to size/location/invasiveness of BCC.
Vismodegib
Brand(s): Erivedge
DOSAGE FORM/ STRENGTH: 150 mg tablet

Note: Patient preference for oral therapy will not be considered

Renewals will be considered where the physician has confirmed that the patient has not experienced disease progression while on Erivedge therapy.

Duration of Approval: 1 Year
ONCOLOGY – SUPPORTIVE MANAGEMENT

Aprepitant
Brand(s): Emend
DOSAGE FORM/ STRENGTH: 80 mg, 125 mg capsule, Tri-pack

Effective September 25, 2014, Emend transitioned to the ODB formulary for reimbursement in patients who meet the Limited Use criteria. Dosage regimens not meeting the LU criteria may be submitted to the EAP for consideration of reimbursement.

Denosumab
Brand(s): Xgeva
DOSAGE FORM/ STRENGTH: 120 mg per vial for subcutaneous injection

For the treatment of bony metastases in patients with hormone refractory prostate cancer.

Xgeva is considered through CCO for those receiving prostate cancer treatment from a cancer clinic.

Hormone refractory prostate cancer is determined using the following criteria:

i) Patient has an elevated PSA level or evidence of progressive bony disease\(^1\), despite castrate serum testosterone levels (Less than 1.7 nmol/L or less than 50 ng/dL)\(^2\).

\(^1\) Progressive bony disease is defined as progressive changes in radionucleotide bone scan or clinical signs of disease progression, such as pathologic fracture or increasing bone pain.

\(^2\) Note: Patients who have undergone orchidectomy do not need to provide a serum testosterone level in the request submission.

Approved Dosing: 120 mg subcutaneously every four (4) weeks

Duration of Approval: 1 Year

Renewals will be considered for patient responding to treatment with Xgeva and who still requires treatment.

Duration of Approval: 1 Year
Filgrastim [Granulocyte colony stimulating factor (G-CSF)]
Brand(s): Neupogen
DOSAGE FORM/ STRENGTH: 300 mcg/mL, 480 mcg /1.6 mL

Effective August 30, 2017, Exceptional Access Program (EAP) requests for Neupogen (filgrastim) will no longer be accepted for any indication.

Patients who have an existing EAP approval for Neupogen can continue to receive Neupogen for the duration of the EAP approval period.

Neupogen and Grastofil are not interchangeable products. As of August 30, 2017, new prescriptions for filgrastim for ODB eligible patients will be dispensed Grastofil, unless it specifies Neupogen with the appropriate LU code. Refer to the Ministry’s e-formulary for a listing of Limited Use (LU) criteria for Neupogen.

Effective December 22, 2016, the subsequent entry biologic (SEB) filgrastim as Grastofil® is funded under the Ontario Drug Benefit (ODB) Program as a general benefit (GB).

Please refer to the e-formulary for funded strengths.

Zoledronic Acid
Brand(s): Zometa Concentrate
DOSAGE FORM/ STRENGTH: 4 mg/ 5 mL Vial

Zoledronic acid as Zometa Concentrate will only be considered for the treatment of bony metastases in those with hormone refractory prostate cancer as well as other cancers through the Exceptional Access Program (EAP) in those receiving outpatient care who do not meet the criteria of Cancer Care Ontario (CCO).

Zometa is considered through CCO for those receiving prostate cancer treatment from a cancer clinic.

Duration of Approval: 6 Months

For the treatment of bony metastases for patients with hormone refractory prostate cancer as determined by an elevated PSA level, or evidence of progressive bony disease¹, despite castrate serum testosterone levels (<50 ng/dL).

¹Progressive bony disease should be demonstrated by: progressive changes in radionucleotide bone scan or clinical signs of disease progression (e.g., via radionucleotide scanning, pathologic fracture or increasing bone pain).

Requests for patients who have undergone orchidectomy do not need to provide a serum testosterone level.
Zoledronic Acid
Brand(s): Zometa Concentrate
DOSAGE FORM/ STRENGTH: 4 mg/ 5 mL Vial

- For the prevention of skeletal related events in patients who have not experienced previous skeletal related events\(^2\) and who have bony metastases secondary to:

  - solid tumours (e.g. renal, small cell lung, pancreatic cancers) who have good performance status\(^3\) OR

  - breast cancer or multiple myeloma who are intolerant to pamidronate.

\(^2\)A skeletal related event is defined as: pathologic fracture, spinal cord compression, radiation therapy to bone or surgery to bone.

\(^3\)Good performance status is defined as patients that are ambulatory, capable of self care and up and about more than 50 per cent of waking hours.

- For the treatment of patients with symptoms due to bony metastases secondary to breast cancer or multiple myeloma who have failed or are intolerant to pamidronate.

- Consideration for patients who are symptomatic due to bony metastases secondary to other types of solid tumours or cancers will be considered on a case-by-case basis. The physician is asked to include information describing the patient’s bone pain and use of other therapies including the use of bisphosphonates. The use of other non-pharmacologic treatment modalities such as surgery or radiation that have been tried should be provided in the request.

**Duration of Approval:** 6 Months

**Renewals** will be considered for patients who are responding to therapy and is still deemed to require treatment.

**Duration of Approval:** 6 Months
OSTEOPOROSIS

Teriparatide
Brand(s): Forteo
DOSAGE FORM/ STRENGTH: 250 mcg/mL - 3 mL prefilled pen, 250 mcg/mL 2.4 mL prefilled pen

For the treatment of osteoporosis in patients who meet the following criteria;

- 65 years of age or older who are mobile; **AND**
- Patient is at high risk of fragility fractures*; **AND**
- Patient who has osteonecrosis of the jaw due to an anti-resorptive agent **OR** who has atypical femur fracture due to an anti-resorptive agent. (Note: One of the two conditions must be present.)

*High risk for fragility fractures is defined as

- A bone mineral density (BMD) T-score less than or equal to -3; **AND**
- Prior fragility fracture

Note: Requesting physicians must include a copy of the BMD report with the EAP request

Requests meeting criteria will be funded for 24 months. It should be noted that renewals are NOT considered.

(As noted in the product monograph, the maximum life time exposure to an individual patient is 24 months)

1No other contraindications to anti-resorptive therapies will be considered for funding.

Duration of Approval: Total approval duration of 24 months will be provided. Renewals are not considered.
PAIN MANAGEMENT

Cannabidiol and delta-9-tetrahydro-cannabinol
Brand(s): Sativex
DOSAGE FORM/ STRENGTH: 25 mg/27 mg per mL buccal spray

For the treatment of **neuropathic pain related to multiple sclerosis** in patients who have:

- Ineffective response or intolerable side effects / contraindications to adequate trials* of a tricyclic antidepressant and gabapentin and pregabalin; and

- Ineffective response or intolerable side effects / contraindications to adequate trials* of Cesamet (nabilone) and Marinol (delta-9-tetrahydrocannabinol); and

- No contraindications to Sativex therapy.

* Adequate trial is defined as 2 months unless intolerable side effect(s) occur.

**Duration of Approval**: 1 Year

Note: Side effects and contraindications must be described in detail. Side effects should be deemed serious by the physician such that no further therapy with the agent would be warranted.

**Renewal** will be considered for patients responding to Sativex therapy as demonstrated by decreased pain and other pain-related symptoms; no initiation of new analgesics; and no increase in doses of any analgesics.

**Duration of Approval**: Renewal is lifetime.

Sativex is also reimbursed for the treatment of refractory pain in palliative cancer patients according to specified criteria.

**Duration of Approval**: 6 Months
**Methadone**
Brand(s): Metadol
DOSAGE FORM/ STRENGTH: 1 mg, 5 mg, 10 mg, 25 mg tablets, 1 mg/mL oral solution, 10 mg/mL oral concentrate solution

For the treatment of cancer and non-cancer pain in patients who cannot tolerate, or have failed treatment with a listed long-acting opioid.

The CED noted that there is a potential for drug interactions with the use of methadone resulting from inhibition of drug metabolism (via CYP 3A4 inhibition; e.g. QT prolongation with certain antibiotics). The requesting physician is asked to ensure that this issue is addressed with the patient.

Duration of Approval 1 Year

**Renewals** will be considered on a case-by-case basis.

For renewals, the requesting physician is asked to provide details of the patient’s clinical response to therapy and additional information pertaining to the current medications and addition or stoppage of other pain medications in the prior year of methadone use. Please specify the dosages and dosing frequency of current medications and provide reasons for any changes in the medication regimen.

**Oxycodone Controlled Release Tablet**
Brand(s): OxyNeo
DOSAGE FORM/ STRENGTH: 10 mg CR, 15 mg CR, 20 mg CR, 30 mg CR, 40 mg CR

For the treatment of chronic pain in patients who have experienced intolerance or have failed an adequate trial (for example, three months) of at least one other listed long-acting opioid product.

**Note**: Physicians should consider best practice guidelines for the safe and effective use of opioids in chronic non-cancer pain, such as the Canadian Guideline for Safe and Effective Use of Opioids for Chronic Non-Cancer Pain.

Please include the following information in your request:

iv) The diagnosis for which the pain management is required must be documented.

v) All concomitant pain medication therapy must be documented.

vi) Other medications with potential for abuse or interaction with opioid therapy should be documented.

**Duration of Approval**: 1 Year
Oxycodone Controlled Release Tablet
Brand(s): OxyNeo
DOSAGE FORM/ STRENGTH: 10 mg CR, 15 mg CR, 20 mg CR, 30 mg CR, 40 mg CR

Renewals will be considered if treatment continues to be appropriate for the management of the patient’s chronic pain. Please include the following information on your renewal request:

Duration of Approval: 1 Year

i) All concomitant pain medication therapy must be documented.

ii) Other medications with potential for abuse or interaction with opioid therapy should be documented.

Note: OxyNEO 60mg and 80mg tablets are not funded.

Note: Physicians registered on the Ontario Medical Association’s Palliative Care Facilitated Access List and Nurse Practitioners registered on the PCFA list through the Registered Nurses Association of Ontario or the Nurse Practitioner Association of Ontario can access OxyNeo for chronic pain management of their palliative care patient for an initial duration of one year without approval through the Exceptional Access Program.
PARKINSON’S DISEASE TREATMENTS

Apomorphine
Brand(s): Movapo
DOSAGE FORM/ STRENGTH: 30mg/3mL Prefilled Multi-dose Pen for Injection

Initiation Criteria:

For the acute, intermittent treatment of patients with advanced Parkinson’s disease (PD) who meet the following criteria;

a) Apomorphine is used as adjunctive treatment in patients receiving optimal PD treatment with a levodopa-based drug and a dopamine agonist (DA); AND

b) Despite receiving optimal treatment, the patient continues to experience intermittent hypomobility defined as either “off” episodes as the drug wears off or unpredictable on/off episodes; AND

b) Patient is under the care of a physician experienced in Parkinson’s disease who is practicing in one of the province’s specialized movement disorder clinics or in consultation with an expert from one of these clinics (consult note must be included).

1 Optimal treatment is defined as receiving maximally tolerated dose of a levodopa based therapy (such as levodopa/carbidopa or levodopa/benserazide) AND a dopamine agonist (such as bromocriptine, pramipexole, ropinirole, rotigotine) for at least 30 days. If the patient is intolerant to or unable take a DA, provide a description of the intolerance or clinical contraindication. In such cases, another class of PD therapy [i.e. the catechol-O-methyl transferase (COMT) inhibitor (such as entacapone) or a monoamine oxidase-B inhibitor (such as rasagiline, selegiline)] may be considered with the levodopa-based regimen.

2 An Ontario specialized movement disorder clinic listed on the website of the Canadian Movement Disorder Group [http://www.cmdg.org/AcrossCanada/acrosscanada.htm#que](http://www.cmdg.org/AcrossCanada/acrosscanada.htm#que) (Patients in border communities seeing specialists in a listed Manitoba or Quebec clinic are also eligible.)

Exclusion:
Concomitant use with 5-hydroxytryptamine type 3 (5HT3) antagonists (e.g. ondansetron, granisetron, dolasetron)

Recommended dose: 0.2 mL (2 mg) to 0.6 mL (6 mg) per dose, to be administered subcutaneously as an adjunct to regular oral anti-PD medications.
Apomorphine Hydrochloride
Brand(s): Movapo
DOSAGE FORM/ STRENGTH: 30mg/3mL Prefilled, Multi-dose Pen for Injection

Renewal criteria:

Renewals will be considered in patients who continue to benefit from treatment and who do not develop unacceptable toxicities to treatment with apomorphine. At the time of renewal, please confirm the dose and average frequency of use.

Provide an improved outcome resulting from treatment with apomorphine (e.g. improvement in the frequency or duration of mobility or hypomobility, duration of off episodes, quality of life measure, symptom improvements) compared to baseline before treatment with apomorphine.

Approval duration of initials and renewals: 1 year

Levodopa 20 mg/mL and Carbidopa 5 mg/mL Intestinal gel
Brand(s): Duodopa
DOSAGE FORM/ STRENGTH: Intestinal Gel containing Levodopa 20 mg/mL – Carbidopa 5 mg/mL (100 mL cassette)

For the treatment of Parkinson's disease in patients who meet the following criteria;

- Experiences at least 25% of the waking day in the off state; AND
- Has severe disability while in the off-state as assessed by a Movement Disorder Specialist; AND
- Has received an adequate trial of maximally tolerated doses of levodopa, with demonstrated clinical response; AND
- Has failed adequate trials of other adjunctive medications (entacapone, dopamine agonists, monoamine oxidase-B [MAO-B] inhibitors) if not contraindicated. Note that if a contraindication is deemed to be applicable to the patient, the requesting physician must state the contraindication and provide the rationale why it is considered a contraindication for the patient).

Clinical details pertaining to the severity of the patient’s disability while in the off-state as well as a complete history of all previous and current medications (e.g., name, start date and duration of therapy, doses used, side effects, and response) must be included.

Requests for treatment initiation will be limited to the physicians practicing in the following specialized movement disorder clinics: Ottawa, London, Toronto Western, Kingston, Baycrest and Hamilton. (Note: An Ontario specialized movement disorder clinic listed on the website of the Canadian Movement Disorder Group http://www.cmdg.org/AcrossCanada/acrosscanada.htm#que is acceptable)
Levodopa 20mg/mL and Carbidopa 5mg/mL Intestinal gel
Brand(s): Duodopa
DOSAGE FORM/ STRENGTH: Intestinal gel containing Levodopa 20mg/mL and Carbidopa 5mg/mL (100 mL cassette)

Exclusion criteria: Patients meeting ANY of the following criteria will NOT be considered.
- Patients who have a contraindication to insertion of a percutaneous endoscopic gastrostomy (PEG) tube
- Severe psychosis or dementia

Duration of Approval of Initials: 1 Year

Renewals will be considered in patients who continue to benefit from treatment. The patient should continue to demonstrate a significant reduction in the time spent in the off state and an improvement in the severity of the disability in the off state.

Duration of Approval of Renewals: 1 Year

Rasagiline
Brand(s): Azilect
DOSAGE FORM/ STRENGTH: 0.5 mg, 1 mg tablet

For the treatment of patients with Parkinson’s disease who experience about 25% of the waking day in the off-state despite maximally tolerated doses of levodopa.

Duration of Approval of Initials and Renewals: 5 Years
PSORIATIC ARTHRITIS TREATMENTS

Adalimumab – See Formulary for funded biosimilars
Brand(s): Humira and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 40 mg/0.8 mL prefilled syringe, 40 mg/0.8mL and 20 mg/0.2 mL prefilled pens for subcutaneous injection

Certolizumab
Brand(s): Cimzia
DOSAGE FORM/ STRENGTH: 200 mg/mL prefilled syringe and autoinjector

Etanercept – see Formulary for funded biosimilars
Brand(s): Enbrel and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 25 mg/vial and 50 mg prefilled syringe or pens for subcutaneous injection per formulary listed options

Golimumab
Brand(s): Simponi
DOSAGE FORM/ STRENGTH: 50 mg/0.5 ml prefilled syringe and autoinjector

Secukinumab
Brand(s): Cosentyx
DOSAGE FORM/ STRENGTH: 150 mg/mL prefilled syringe and 150 mg/mL prefilled pen

For the treatment of psoriatic arthritis in patients who have:
Severe active disease (≥ 5 swollen joints and radiographic evidence of psoriatic arthritis) despite treatment with methotrexate (20mg/week) for at least 3 months and one of leflunomide (20 mg/day) or sulfasalazine (1 g twice daily) for at least 3 months.

If the patient has documented contraindications or intolerances to methotrexate, then only one of leflunomide (20 mg/day) or sulfasalazine (1 g twice daily) for at least 3 months is required. Details of contraindications and intolerances must also be provided.

Duration of Approval of initials: 1 Year

Renewal will be considered for patients with objective evidence of at least a 20% reduction in swollen joint count and a minimum of improvement in 2 swollen joints over the previous year. For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided.

Duration of Approval of first renewal: 1 Year
The planned dosing regimen for the requested biologic should be provided. The recommended doses for the treatment of psoriatic arthritis are as follows:

- Adalimumab 40mg every two weeks
- Certolizumab 400 mg at week 0, 2, 4 then maintenance doses of 200 mg every 2 weeks or 400 mg every 4 weeks
- Etanercept 25 mg twice weekly or 50mg once weekly
- Golimumab 50 mg once a month
- Secukinumab 150mg sc at weeks 0, 1, 2, and 3 followed by monthly maintenance dosing starting at week 4. If a patient is an anti-TNFalpha inadequate responder and continues to have active psoriatic arthritis, consider using the 300 mg sc dose.

For psoriatic arthritis patients with coexistent moderate to severe plaque psoriasis, use the dosing and administration recommendations for plaque psoriasis (i.e. 300 mg sc at weeks 0, 1, 2, and 3, followed by monthly maintenance dosing starting at week 4)

**Duration of Approval of second and subsequent renewals:** 5 years
For the treatment of psoriatic arthritis (PsA) in patients who have:

Severe active disease (≥ 5 swollen joints and radiographic evidence of psoriatic arthritis) despite treatment with methotrexate (20mg/week) for at least 3 months and one of leflunomide (20mg/day) or sulfasalazine (1g twice daily) for at least 3 months.

If the patient has documented contraindications or intolerances to methotrexate, then only one of leflunomide (20mg/day) or sulfasalazine (1g twice daily) for at least 3 months is required. Details of contraindications and intolerances must also be provided.

**Duration of Approval**: 1 Year

**Renewals** will be considered for patients with objective evidence of at least a 20% reduction in swollen joint count and a minimum of improvement in 2 swollen joints over the previous year.

For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided.

**Duration of Approval of first renewal**: 1 Year

**Duration of Approval**: Second and subsequent renewals are 2 years

The planned dosing regimen for the requested biologic should be provided.

**Recommended Dose**:

For psoriatic arthritis (PsA) patients or those with PsA and coexistent mild plaque psoriasis: 160 mg (two 80 mg injections) sc at Week 0, followed by 80 mg every 4 weeks.

For psoriatic arthritis patients with coexistent moderate-to-severe plaque psoriasis, you may wish to refer to the ODB formulary for access upon meeting the Limited Use Criteria for Plaque psoriasis. (EAP authorization would not be required).

Dose recommended for such patients is 160 mg by subcutaneous injection (two 80 mg injections) at Week 0, followed by 80 mg every 2 weeks for 6 doses (i.e. weeks 2, 4, 6, 8, 10, and 12), then 80 mg every 4 weeks thereafter.

Ixekizumab may be used alone or in combination with a conventional DMARD (e.g. methotrexate)
PULMONARY ARTERIAL HYPERTENSION

Drugs for Pulmonary Arterial Hypertension (PAH) under EAP

i) Phosphodiesterase (PDE)-5 inhibitor: sildenafil (Revatio), tadalafil (Adcirca)

ii) Endothelin receptor antagonists (ERAs): ambrisentan (Volibris), bosentan (Tracleer), macitentan (Opsumit)

iii) Prostanoids: epoprostenol (Flolan, Caripul), treprostinil (Remodulin), selexipag (Uptravi)

Sildenafil
Brand(s): Revatio
DOSAGE FORM/ STRENGTH: 20 mg tablet

Tadalafil
Brand(s): Adcirca
DOSAGE FORM/ STRENGTH: 20 mg tablet

All requests (initial, renewal, monotherapy, combination therapy) for a PAH drug must come from one of the following recognized PAH referral centres.

- Pulmonary Hypertension Centre
  Hamilton Health Sciences – General Hospital
- The Firestone Institute Pulmonary Hypertension Program
  St. Joseph's Healthcare Hamilton and McMaster University
- Pulmonary Hypertension Clinic
  Hotel Dieu Hospital/Kingston General Hospital
- Pulmonary Hypertension Program
  London Health Science Centre – Victoria Hospital
- Ottawa Pulmonary Hypertension Clinic
  University of Ottawa Heart Institute and the Ottawa Hospital
- University Health Network Pulmonary Hypertension Program
  Toronto General Hospital
Requests from other physicians/centres must include a recent (less than or equal to 3 months old) consult note/recommendation from a recognized PAH referral centre that supports the request;

Out-of-province referral centre consults (e.g., from Winnipeg for patients in Northern Ontario) will also be considered on a case-by-case basis

i) **Sildenafil** (Revatio, generics), **Tadalafil** (Adcirca, generics)

Initial Criteria:

For the treatment of patients with pulmonary arterial hypertension (PAH) [WHO Group 1 Pulmonary hypertension] who meet all the following criteria;

- PAH defined as a resting mean pulmonary artery pressure (mPAP) of ≥ 25 mmHg at rest AND normal pulmonary capillary wedge pressure (PCWP) ≤ 15 mmHg on right heart catheterization\(^1\); AND

- The drug request meets one of the following circumstances of use:
  - Drug is being used as monotherapy in a patient with WHO-functional class II (Note that a PDE-5 inhibitor must be used as first line monotherapy for WHO-FC II (unless contraindicated or demonstrated intolerance), III, or IV; OR
  - Drug is being used as sequential dual therapy in combination with a funded ERA (i.e. ambrisentan, bosentan, macitentan) or a funded prostanoid (i.e. epoprostenol, treprostinil) in a patient who has had an inadequate response with monotherapy (i.e., failure to achieve WHO-FC I or II; or 6MWD >440 metres; or no/mild RV failure); OR
  - Drug is being used as up-front dual therapy in combination with a funded ERA (i.e. ambrisentan, bosentan, macitentan) or a funded prostanoid (i.e. epoprostenol, treprostinil) in a patient with advanced disease (i.e. WHO-functional class III or IV, 6MWD <380 metres; OR evidence of RV failure.)

\(^1\) Note: Left ventricular end-diastolic pressure ≤ 15 mmHg is also acceptable.
Ambrisentan
Brand(s): Volibris
DOSAGE FORM/ STRENGTH: 5 mg, 10 mg tablet

Bosentan
Brand(s): Tracleer, Generics (Co-, Mylan-, PMS-, Sandoz-)
DOSAGE FORM/ STRENGTH: 62.5 mg, 125 mg tablet

Macitentan
Brand(s): Opsumit
DOSAGE FORM/ STRENGTH: 10 mg tablet

Updated April 20, 2021

All requests (initial, renewal, monotherapy, combination therapy) for a PAH drug must come from one of the following recognized PAH referral centres.

- **Pulmonary Hypertension Centre**
  Hamilton Health Sciences – General Hospital

- **The Firestone Institute Pulmonary Hypertension Program**
  St. Joseph's Healthcare Hamilton and McMaster University

- **Pulmonary Hypertension Clinic**
  Hotel Dieu Hospital/Kingston General Hospital

- **Pulmonary Hypertension Program**
  London Health Science Centre – Victoria Hospital

- **Ottawa Pulmonary Hypertension Clinic**
  University of Ottawa Heart Institute and the Ottawa Hospital

- **University Health Network Pulmonary Hypertension Program**
  Toronto General Hospital

Requests from other physicians/centres must include a recent (less than or equal to 3 months old) consult note/recommendation from a recognized PAH referral centre that supports the request;

Out-of-province referral centre consults (e.g., from Winnipeg for patients in Northern Ontario) will also be considered on a case-by-case basis

Initial Criteria:

For the treatment of patients with pulmonary arterial hypertension (PAH) [WHO Group 1 Pulmonary hypertension] who meet all the following criteria;
Ambrisentan  
Brand(s): Volibris  
DOSAGE FORM/ STRENGTH: 5 mg, 10 mg tablet

Bosentan  
Brand(s): Tracleer, Generics (Co-, Mylan-, PMS-, Sandoz-)  
DOSAGE FORM/ STRENGTH: 62.5 mg, 125 mg tablet

Macitentan  
Brand(s): Opsumit  
DOSAGE FORM/ STRENGTH: 10 mg tablet  
Updated: April 20, 2021

- PAH defined as a resting mean pulmonary artery pressure (mPAP) of $\geq 25 \text{ mmHg}$ at rest AND normal pulmonary capillary wedge pressure (PCWP) $\leq 15 \text{ mmHg}$ on right heart catheterization\(^1\); AND

- The drug request meets one of the following circumstances of use:
  - Drug is being used as monotherapy in a patient with WHO-functional class III or IV; OR
  - Drug is being used as monotherapy in a patient with WHO-functional class II who has contraindication or has intolerance to a PDE-5 inhibitor; OR
  - Drug is being used as sequential dual therapy in combination with a funded PDE-5 (i.e. sildenafil, tadalafil) or a funded prostanoid (i.e. epoprostenol, treprostinil) in a patient who has had an inadequate response with monotherapy (i.e., failure to achieve WHO-FC I or II; or 6MWD >440 metres; or no/mild RV failure); OR
  - Drug is being used as up-front dual therapy in combination with a funded PDE-5 (i.e. sildenafil, tadalafil) or a funded prostanoid (i.e. epoprostenol, treprostinil) in a patient with advanced disease (i.e. WHO-functional class III or IV; OR 6MWD <380 metres; OR evidence of RV failure.)

\(^1\) Note: Left ventricular end-diastolic pressure $\leq 15 \text{ mmHg}$ is also acceptable.
Treprostinil
Brand(s): Remodulin
DOSAGE FORM/ STRENGTH: 1 mg/mL, 2.5 mg/mL, 5 mg/mL and 10 mg/mL vials

Epoprostenol
Brand(s): Flolan
DOSAGE FORM/ STRENGTH: 0.5 mg and 1 mg vial

Epoprostenol
Brand(s): Caripul
DOSAGE FORM/ STRENGTH: 0.5 mg and 1.5 mg vial

All requests (initial, renewal, monotherapy, combination therapy) for a PAH drug must come from one of the following recognized PAH referral centres.

- **Pulmonary Hypertension Centre**
  Hamilton Health Sciences – General Hospital
- **The Firestone Institute Pulmonary Hypertension Program**
  St. Joseph's Healthcare Hamilton and McMaster University
- **Pulmonary Hypertension Clinic**
  Hotel Dieu Hospital/Kingston General Hospital
- **Pulmonary Hypertension Program**
  London Health Science Centre – Victoria Hospital
- **Ottawa Pulmonary Hypertension Clinic**
  University of Ottawa Heart Institute and the Ottawa Hospital
- **University Health Network Pulmonary Hypertension Program**
  Toronto General Hospital

Requests from other physicians/centres must include a recent (less than or equal to 3 months old) consult note/recommendation from a recognized PAH referral centre that supports the request;

Out-of-province referral centre consults (e.g., from Winnipeg for patients in Northern Ontario) will also be considered on a case-by-case basis.
Treprostinil
Brand(s): Remodulin
DOSAGE FORM/ STRENGTH: 1 mg/mL, 2.5 mg/mL, 5 mg/mL and 10 mg/mL vials

Epoprostenol
Brand(s): Flolan
DOSAGE FORM/ STRENGTH: 0.5 mg and 1mg vial

Epoprostenol
Brand(s): Caripul
DOSAGE FORM/ STRENGTH: 0.5 mg and 1.5 mg vial

Epoprostenol (Flolan, Caripul), Treprostinil (Remodulin)

Initial Criteria:

For the treatment of patients with pulmonary arterial hypertension (PAH) [WHO Group 1 Pulmonary hypertension] who meet all the following criteria;

- PAH defined as a resting mean pulmonary artery pressure (mPAP) of \( \geq 25 \) mmHg at rest AND normal pulmonary capillary wedge pressure (PCWP) \( \leq 15 \) mmHg on right heart catheterization\(^{1}\); AND

- The drug request meets one of the following circumstances of use:
  - Drug is being used as monotherapy in a patient with WHO-functional class III or IV; OR
  - Drug is being used as sequential dual therapy in combination with a funded PDE-5 (i.e. sildenafil, tadalafil) or with a funded ERA (i.e. ambrisentan, bosentan, macitentan) in a patient who fails to meet treatment targets (i.e. failure to achieve WHO-FC I or II; or 6MWD >440 metres; or no/mild RV failure) with monotherapy; OR
  - Drug is being used as up-front dual therapy in combination with a funded PDE-5 (i.e. sildenafil, tadalafil) or with a funded ERA (i.e. ambrisentan, bosentan, macitentan) in a patient with advanced disease (i.e. WHO-functional class III or IV; OR 6MWD <380 metres; OR evidence of RV failure.)

\(^{1}\) Note: Left ventricular end-diastolic pressure \( \leq 15 \) mmHg is also acceptable.
Treprostinil  
Brand(s): Remodulin  
DOSAGE FORM/ STRENGTH: 1 mg/mL, 2.5 mg/mL, 5 mg/mL and 10 mg/mL vials  

Epoprostenol  
Brand(s): Flolan  
DOSAGE FORM/ STRENGTH: 0.5 mg and 1mg vial  

Epoprostenol  
Brand(s): Caripul  
DOSAGE FORM/ STRENGTH: 0.5 mg and 1.5 mg vial  

For all funded PAH Drugs, case-by-case consideration may be provided for the following:  

- Requests for triple therapy (Including patients awaiting lung transplant.)  
- Patients who may have mixed co-morbidities that include ILD, COPD or LV failure. (i.e. patients with mixed WHO Group 1 and Group 3 pulmonary hypertension OR mixed WHO Group 1 and Group 2 pulmonary hypertension)  

Exclusion Criteria:  
Combinations of drugs targeting similar pathways will not be funded. (i.e. combination regimen may only include one agent from each drug class -- phosphodiesterase type 5 [PDE-5] inhibitors, endothelin receptor antagonists (ERA), and/or prostanoids)  

Renewal criteria for funded PAH Drugs:  
Renewals will be provided for patients who remain under the care of a physician from a recognized PAH Centre (see list above) and who continue to benefit from therapy.  

Approval Durations:  
Duration of Approval for Initial Requests: 1 year  
Duration on triple therapy regimens awaiting lung transplantation: 1 year  
Duration of first renewal: 1 Year  
Duration of subsequent renewals: 5 Years
All requests (initial, renewal, monotherapy, combination therapy) for a PAH drug must come from one of the following recognized PAH referral centres.

- **Pulmonary Hypertension Centre**
  Hamilton Health Sciences – General Hospital
- **The Firestone Institute Pulmonary Hypertension Program**
  St. Joseph's Healthcare Hamilton and McMaster University
- **Pulmonary Hypertension Clinic**
  Hotel Dieu Hospital/Kingston General Hospital
- **Pulmonary Hypertension Program**
  London Health Science Centre – Victoria Hospital
- **Ottawa Pulmonary Hypertension Clinic**
  University of Ottawa Heart Institute and the Ottawa Hospital
- **University Health Network Pulmonary Hypertension Program**
  Toronto General Hospital

**Initial Criteria:**

For the treatment of patients with pulmonary arterial hypertension (PAH) [WHO Group 1 Pulmonary hypertension] who meet all the following criteria:

- PAH defined as a resting mean pulmonary artery pressure (mPAP) of ≥ 25 mmHg at rest AND normal pulmonary capillary wedge pressure (PCWP) ≤ 15 mmHg on right heart catheterization; AND
- Patient with World Health Organization (WHO) functional class II to IV; AND
- Selexipag is being used in a patient experiencing inadequate control with a Phosphodiesterase (PDE)-5 inhibitor (i.e. tadalafil or sildenafil) AND an endothelin receptor antagonist (ERA) (i.e. bosentan, ambrisentan, or macitentan)

**Notes:**

1. Left ventricular end-diastolic pressure ≤ 15 mmHg is also acceptable.
2. Unable to meet treatment targets (i.e. failure to achieve WHO-FC I or II; or 6MWD >440 metres; or no/mild RV failure)

Case-by-case consideration may be provided for the following;

- Requests for Selexipag in patients who demonstrate intolerance or have a contraindication to either PDE-5 inhibitors (i.e. both sildenafil and tadalafil) or ERAs (i.e. each of bosentan, ambrisentan, macitentan)
Selexipag
Brand(s): Uptravi
DOSAGE FORM/ STRENGTH: 200mcg, 400mcg, 600mcg, 800mcg, 1000mcg, 1200mcg, 1400mcg, 1600mcg Tablets

- Patients who may have mixed co-morbidities that include ILD, COPD or LV failure. (i.e. patients with mixed WHO Group 1 and Group 3 pulmonary hypertension OR mixed WHO Group 1 and Group 2 pulmonary hypertension)

Exclusion Criteria:
Combination therapy with prostacyclin or prostacyclin analog therapies and Selexipag will not be covered.

Renewal criteria:
Renews will be provided for patients who remain under the care of a physician from a recognized PAH Centre (see list above) and who continue to benefit from therapy.

Approval Durations:
Duration of Approval for Initial Requests: 1 year
Duration on triple therapy regimens awaiting lung transplantation: 1 year
Duration of first renewal: 1 Year
Duration of subsequent renewals: 5 Years
Riociguat
Brand(s): Adempas
DOSAGE FORM/ STRENGTH: 0.5 mg, 1 mg, 1.5 mg, 2 mg, 2.5 mg tablet

For the treatment of chronic thromboembolic pulmonary hypertension (CTEPH) in patients who meet the following criteria;

- the physician making the request is a clinician with experience in the diagnosis and treatment of CTEPH\(^1\); AND

- the patient is diagnosed with inoperable CTEPH (World Health Organization [WHO] Group 4); OR persistent or recurrent CTEPH after surgical treatment in adult patients (18 years of age or older) with WHO Functional Class (FC) II or III pulmonary hypertension.

\(^1\) Request should come from a clinician from a Pulmonary Hypertension referral centre (See Pulmonary Arterial Hypertension referral clinics above).

**Duration of Approval:** 1 Year

**Renewal** of funding will be considered for patients who continue to respond to therapy with riociguat. When submitting a request for renewal of funding, the physician should submit clinical information to support that the patient is deriving benefit from the treatment compared to before they started the treatment. The physician should provide confirmation of improvement of any ONE or more reasonable clinical parameters which supports the response of the patient’s CTEPH to riociguat.

**Duration of Approval:** 1 Year

**Requests for subsequent funding renewals** (i.e. beyond the first two years of treatment) will be considered when a physician provides written confirmation that the patient continues to respond to therapy with riociguat. The physician should provide confirmation of improvement of any ONE or more reasonable clinical parameters which supports the response of the patient’s CTEPH to riociguat compared to baseline or that supports that the patient’s condition is stable while on riociguat.

**Duration of Approval:** Subsequent Renewals - 5 Years
For the treatment of cystic fibrosis (CF) in patients who meet all of the following criteria:
  - 12 years of age and older; **AND**
  - Confirmed diagnoses of CF with at least one F508del mutation in the cystic fibrosis transmembrane conductance regulator (CFTR gene); **AND**
  - Patient has been optimized on best supportive care for their CF prior to starting Trikafta; **AND**
  - Has a baseline percent predicted forced expiratory volume in 1 second (ppFEV\(_1\)) that is less than or equal to 90%\(^1\) before starting Trikafta; **AND**
  - Prescribed by a clinical specialist affiliated with a Canadian cystic fibrosis centre.

\(^1\)Case-by-case consideration may be provided for patients not meeting lung function criteria.

The following measurements must be completed prior to initiating treatment with Trikafta:

  - Baseline spirometry measurements of FEV\(_1\) in litres and percent predicted (within the last 3 months); **AND**
  - Number of days treated with oral and/or intravenous (IV) antibiotics for pulmonary exacerbations in the previous 6 months OR number of pulmonary exacerbations requiring oral and/or IV antibiotics in the previous 6 months; **AND**
  - Number of CF-related hospitalizations in the previous 6 months; **AND**
  - Weight, height, and body mass index (BMI); **AND**
  - CFQ-R Respiratory Domain score.

**Exclusion criteria for Initial and Renewal criteria:**
(Patients meeting any of the following will not be funded)

  - Patient has undergone lung transplantation; OR
  - Patient is using Trikafta as combination therapy with another cystic fibrosis transmembrane conductance regulator (CFTR) modulator.

**Initial approval duration:** 7 months
Elexacaftor/Tezacaftor/Ivacaftor and Ivacaftor
Brand(s): Trikafta
DOSAGE FORM/ STRENGTH: 100 mg/50 mg/75 mg and 150 mg

Initial Renewal Criteria:
Renewal of funding will be considered in patients demonstrating at least ONE of the following improvements after 6 months of treatment with Trikafta;

1. Improvement of percent predicted FEV₁ by 5% or more above the baseline measurement; OR
2. A decrease in the total number of days for which the patient received treatment with oral and/or IV antibiotics for pulmonary exacerbations compared with the 6-month period prior to initiating treatment OR a decrease in the total number of pulmonary exacerbations requiring oral and/or IV antibiotics compared with the 6-month period prior to initiating treatment; OR
3. Decreased number of CF-related hospitalizations in the 6 months after initiation of Trikafta treatment compared with the 6-month period prior to initiating Trikafta; OR
4. No decline in BMI at 6 months compared with the baseline BMI assessment; OR
5. Improvement by 4 points or more in the CFQ-R Respiratory Domain scale compared to baseline scores.

Subsequent renewal criteria:
For patients who have met the initiation criteria and initial renewal criteria.

Ongoing renewal of funding will be provided for those who are continuing to benefit from therapy with Trikafta and who do not meet any of the exclusion criteria.

At the time of renewal application, please include the patient’s most recent ppFEV₁ and a clinical update to confirm the treatment benefits or response experienced by the patient.

Approval Duration of renewals: 1 year

Approved doses: 12 years of age and older: 2 tablets (each containing elexacaftor/ tezacaftor/ ivacaftor 100mg/ 50mg/ 75mg) taken in the morning & one tablet (ivacaftor 150mg) taken in the evening approximately 12 hours apart.
Ivacaftor  
Brand(s): Kalydeco  
DOSAGE FORM/ STRENGTH: 150 mg tablets  
Effective date: June 20, 2014, Updated December 20, 2019

For the treatment of cystic fibrosis in Patients meeting the following criteria;

(i) the Patient is at least 6 years old and has one of the following mutations in the Cystic Fibrosis Transmembrane conductance Regulator (CFTR) gene: G551D, G1244E, G1349D, G178R, G551S, S1251N, S1255P, S549N or S549R;  
OR  
the Patient is at least 18 years old with an R117H mutation in the CFTR gene.

Initial approval period: 1 year

Initial renewal criteria:

Documented response to treatment (after at least 6 months of therapy), as evidenced by the following:

(a) In cases where the patient’s sweat chloride levels prior to commencing therapy were above 60mmol/litre:
   • the Patient’s sweat chloride level fell below 60mmol/litre; or  
   • the Patient’s sweat chloride level is 30% lower than the level reported in a previous test;
(b) In cases where the patient’s sweat chloride levels prior to commencing therapy were below 60mmol/litre:
   • the Patient’s sweat chloride level is 30% lower than the level reported in a previous test; or  
   • the patient demonstrates a sustained absolute improvement in FEV1 of at least 5% when compared to the FEV1 test conducted prior to the commencement of therapy.

Duration of approval: 1 year

Subsequent renewal criteria: The Patient is continuing to benefit from therapy with Kalydeco.

1It should be noted that, while baseline sweat chloride levels and FEV1 are not required to meet initial approval criteria for Kalydeco, these parameters are used to evaluate the effect of Kalydeco at the time of renewal. To avoid delays, the prescriber should submit a copy of the mutation report, recent baseline sweat chloride levels before starting Kalydeco, and recent baseline FEV1 with the initial request for funding of Kalydeco. These baseline values will be used to evaluate the patient’s response to therapy at the time of renewal and would be logistically difficult to obtain once treatment is initiated.

Duration of Approval: 1 Year
Ivacaftor/Lumacaftor
Brand(s): Orkambi
DOSAGE FORM/ STRENGTH: 100 mg / 125 mg tablets; 200 mg/ 125 mg tablets
Updated: June 25, 2021

Initiation Criteria:

1. Patient is 2 years of age or older: AND
2. Has a confirmed diagnosis of cystic fibrosis and homozygous for F508del mutation in the cystic fibrosis transmembrane conductance regulator gene; AND
3. The patient has demonstrated adherence to their prescribed cystic fibrosis therapeutic regimen; AND
4. Prescribed by a clinical specialist affiliated with a Canadian cystic fibrosis centre: AND
5. Patient must meet one of the following criteria:
   i) Patient is 2 years of age or older and has experienced 1 or more pulmonary exacerbation(s) per year requiring therapy with IV antibiotics OR 3 or more pulmonary exacerbations per year requiring therapy with oral antibiotics.
   ii) Patient is 6 to 11 years of age and has an absolute decline in FEV1 percent predicted equal to or greater than 5% within a 12 month period, sustained over at least 4 months, in spite of optimized medical therapies (For example drop from 90% to 85% predicted)
   iii) Patient is 12 years of age or older and has a baseline FEV1 equal to or less than 70% predicted, who have an absolute decline in FEV1 of ≥5%, within a 12 month period, sustained over at least 4 months, in spite of optimized medical therapies, (For example, FEV1 decline from 60% predicted to 55% predicted in the last 6 months)
   iv) Patient is 12 years of age and older and has a baseline FEV1 of greater than 70% predicted who have an absolute decline in FEV1 of ≥10% predicted within a 12 month period, sustained over at least 4 months, in spite of optimized medical therapies (For example, FEV1 decline from 80% predicted to 70% predicted in the last 6 months)

Exclusion criteria:

(Patients meeting the following will not be funded)

- Patient is currently receiving invasive mechanical ventilation via endotracheal tube or tracheostomy tube; OR
- Patient is the previous recipient of a double lung transplant; OR
- Patient receiving concomitant therapy with another cystic fibrosis transmembrane conductance regulator (CFTR) modulator.
Ivacaftor/Lumacaftor
Brand(s): Orkambi
DOSAGE FORM/ STRENGTH: 100 mg/125 mg tablets; 200 mg/125 mg tablets

Notes:

The following assessments should be made prior to initiating treatment:
- Weight, height, and BMI;
- Number of CF related hospitalizations in the previous 6 months;
- Number of days treated with oral and IV antibiotics for pulmonary exacerbations in the previous 6 months (OR number of pulmonary exacerbations requiring oral and IV antibiotics in previous 6 months);

Additionally, for patients 6 years of age and older all the following MUST be provided:
- Baseline measurement of FEV1 in litres and % predicted taken within 3 months of planned initiation of treatment or prior to commencing treatment;
- Change in FEV1 demonstrating decline in FEV1 % predicted prior to starting therapy (as defined in initiation criteria);
- Cystic Fibrosis Questionnaire Child (CFQ-C) and Cystic Fibrosis Questionnaire-Parent (CFQ-P) for those 6 to 11 years old OR Cystic Fibrosis Questionnaire Respiratory (CFQ-R) Domain score for those 12 years and older.

Approval duration of initials: 7 months

Initial Renewal Criteria:

Renewal of funding will be considered in individuals meeting the following:

1. The patient continues to demonstrate adherence to their prescribed cystic fibrosis therapeutic regimen; AND
2. The patient has demonstrated at least ONE of the following after six months of treatment:
   i) A decrease in the total number of days for which the patient received treatment with oral and/or IV antibiotics for pulmonary exacerbations compared with the six month period prior to initiating treatment OR a decrease in the total number of pulmonary exacerbations requiring oral and IV antibiotics compared with the six month period prior to initiating treatment; OR
   ii) Decreased number of CF related hospitalizations at 6 months compared with the six month period prior to initiating Orkambi treatment; OR
   iii) No decline in BMI at six months compared with the baseline BMI assessment (those 2 to 5 years old may also use BMI percentile); OR
   iv) No decline in FEV1 at six months compared with the baseline FEV1 assessment (as applicable to initiation criteria);
Ivacaftor/Lumacaftor
Brand(s): Orkambi
DOSAGE FORM/ STRENGTH: 100 mg/125 mg tablets; 200 mg/125 mg tablets

3. **If the patient is 6 years of age or older**, they must demonstrate improved or sustained quality of life through an age appropriate test (i.e. CFQ-C and CFQ-P scores for those 6 to 11 years old and CFQ-R Respiratory Domain score if the patient is 12 years or older)

**Subsequent renewal criteria:**

For patients who have met the initiation criteria and initial renewal criteria.

Ongoing renewal of funding will be provided for those who are continuing to benefit from therapy with Orkambi.

**Approval Duration of renewals:** 1 year

**Approved Doses:**

Patients 2 to 5 years old weighing less than 14 kg: lumacaftor 100 mg and ivacaftor 125* mg every 12 hours.

2 to 5 years old weighing more than 14 kg: lumacaftor 150 mg and ivacaftor 188 mg* every 12 hours

6-11 years old: 200 mg/250mg lumacaftor/ivacaftor every 12 hours.

12 years and older: 400 mg/250mg lumacaftor/ivacaftor every 12 hours

*Sachets are currently funded but may be considered case-by-case to accommodate dosing requirements.
Initial approval criteria:

For the treatment of adult patients with a diagnosis of **mild to moderate idiopathic pulmonary fibrosis (IPF)**:

- Diagnosis confirmed by a respirologist and a high-resolution CT scan.
- All other causes of restrictive lung disease (e.g. collagen vascular disorder or hypersensitivity pneumonitis) should be excluded.
- Mild to moderate IPF is defined as forced vital capacity (FVC) greater than or equal to 50% of predicted.
- 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)

**Initial renewal criteria (at 6 months):**

Patients must NOT demonstrate progression of disease defined as an absolute decline in percent predicted FVC of ≥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.

Initial Renewal Duration: 6 Months

**Second and subsequent renewals (at 12 months and thereafter):**

Patients must NOT demonstrate progression of disease defined as an absolute decline in percent predicted FVC of ≥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

**Documentation/information required:**

- **If high-resolution CT scan is not available, lung biopsy may be provided to support the diagnosis of IPF as applicable and available**
- **Full pulmonary function test results.**

Second Renewal Duration: 12 Months

**Exclusion Criteria:**

- Combination use of Ofev (nintedanib) and Esbriet (pirfenidone) will not be funded.
Pirfenidone
Brand(s): Esbriet and generics (see formulary for OFIs)
DOSAGE FORM/STRENGTH: 267 mg capsule, 267 mg tablet, 801 mg tablet

Initial approval criteria:

For the treatment of adult patients with a diagnosis of **mild to moderate idiopathic pulmonary fibrosis (IPF)**:

- Diagnosis confirmed by a respirologist and a high-resolution CT scan.
- All other causes of restrictive lung disease (e.g., collagen vascular disorder or hypersensitivity pneumonitis) should be excluded.
- Mild to moderate IPF is defined as forced vital capacity (FVC) greater than or equal to 50% of predicted.
- 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)

Initial renewal criteria (at 6 months):

Patients must NOT demonstrate progression of disease defined as an absolute decline in percent predicted FVC of ≥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 renewals (at 12 months and thereafter):**

Patients must NOT demonstrate progression of disease defined as an absolute decline in percent predicted FVC of ≥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

**Documentation/information required:**

- If high-resolution CT scan is not available, lung biopsy may be provided to support the diagnosis of IPF as applicable and available
- Full pulmonary function test results.

**Exclusion Criteria:**

Combination use of Esbriet (pirfenidone) and Ofev (nintedanib) will not be funded.
POLYARTICULAR JUVENILE IDIOPATHIC ARTHRITIS

**Etanercept** – see Formulary for funded biosimilars
Brand(s): Enbrel
DOSAGE FORM/ STRENGTH: 25 mg/vial, 25 mg and 50 mg prefilled syringe or pens for subcutaneous injection per formulary listed options

**Adalimumab** – see Formulary for funded biosimilars
Brand(s): Humira and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 40 mg/0.8mL prefilled syringe, 40 mg/0.8mL and and 20 mg/0.2 mL prefilled pens for subcutaneous injection

**Tocilizumab**
Brand(s): Actemra
DOSAGE FORM/ STRENGTH: 80 mg/4 mL Vial, 200 mg/10 mL Vial, 400 mg/20 mL Vial, 162mg/0.9mL Inj (Prefilled syringe), 162mg/0.9mL Auto Injector

Patients with pJIA who are unable to use Erelzi or Brenzys to accommodate weight-based dosing as a result of the syringe format and ability to measure partial syringe doses may request an exemption for Enbrel.

For the first-line treatment of polyarticular-course juvenile idiopathic arthritis in patients meeting the following criteria:

- Patient has active disease (≥ 3 swollen joints and ≥ 5 active joints) despite a trial of optimal dose of subcutaneously administered methotrexate (i.e. 15 mg/m² per week) for at least 3 months. If the patient is unable to tolerate or has a contraindication to subcutaneous methotrexate, the nature of the intolerance or contraindication must be described in detail.

**Duration of Approval:** 1 Year

**Renewal** will be considered for patients with objective evidence of at least a 20% reduction in swollen joint count and a minimum of improvement in 2 swollen joints over the previous year. For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided.

**Duration of Approval:** 5 Year
**Etanercept** – see Formulary for funded biosimilars  
Brand(s): Enbrel  
DOSAGE FORM/ STRENGTH: 25 mg/vial, 25mg or 50 mg prefilled syringe and pens for subcutaneous injection per formulary listed options

**Adalimumab** - see Formulary for funded biosimilars  
Brand(s): Humira and formulary listed biosimilars  
DOSAGE FORM/ STRENGTH: 40 mg/0.8mL prefilled syringe, 40 mg/0.8mL and 20 mg/0.2 mL prefilled pens for subcutaneous injection

**Tocilizumab**  
Brand(s): Actemra  
DOSAGE FORM/ STRENGTH: 80 mg/4 mL Vial, 200 mg/10 mL Vial, 400 mg/20 mL Vial, 162mg/0.9mL Inj (Prefilled syringe), 162mg/0.9mL Auto Injector.

**Dosing for Etanercept:**  
The planned dosing regimen should be provided. The maximum recommended dose is 50mg once weekly.

**Recommended Dosing for Adalimumab:**  
- a) 24 mg/m² (maximum 40 mg) every two weeks; OR
- b) 20 mg every 2 weeks, if the Patient weighs less than 30 kg; OR
- c) 40 mg every 2 weeks, if the Patient weighs more than 30 kg.

**Recommended dosing for tocilizumab in combination with methotrexate:**  

**IV dosing regimen:**  
- a) 10 mg/kg every 4 weeks, if the Patient weighs less than 30kg; OR
- b) 8 mg/kg every 4 weeks, if the Patient weighs more than or equal to 30kg.

**SC dosing regimen:**  
- a) 162 mg once every 3 weeks if the Patient weighs less than 30kg
- b) 162 mg once every 2 weeks if the Patient weighs 30kg or more
Abatacept
Brand(s): Orencia
DOSAGE FORM/ STRENGTH: 250 mg/15 mL vial (Note that the sc injection is not approved for this indication)

Infliximab - See formulary for funded biosimilars
Brand(s): Avsola, Inflectra, Renflexis Biosimilars); Remicade (Originator)
DOSAGE FORM/ STRENGTH: 100 mg/vial

For the treatment of polyarticular-course juvenile idiopathic arthritis in patients meeting the following criteria;

- Patient has active disease (a minimum of 3 (three) swollen joints and a total of 5 active joints); AND

- Patient has had an inadequate response to a three month course of methotrexate administered subcutaneously at a dosage of at least 15 mg/m² per week for at least 3 months. If the patient is unable to tolerate or has a contraindication to subcutaneous methotrexate the nature of the intolerance or contraindication must be described in detail.; AND

- Patient has had an inadequate response to a three month course of etanercept OR adalimumab OR tociluzumab. If the patient is unable to tolerate or has a contraindication to etanercept OR adalimumab OR tociluzumab, the nature of the intolerance or contraindication must be described in detail.

**Duration of Approval:** 1 Year

**Renewals** will be considered for patients with objective evidence of at least a 20% reduction in swollen joint count. For renewals beyond the second year, objective evidence of preservation of treatment effect should be provided. (i.e. the current joint count should be compared to the count prior to initiating treatment with the biologic agent)

**Duration of Approval:** 5 Year

**Approved Dose:**

**Abatacept** refer to the Orencia product monograph for dosing information

**Infliximab** dose up to 6mg/kg/dose at 0, 2 and 6 weeks followed by maintenance of up to 6mg/kg/dose every 8 weeks
RHEUMATOID ARTHRITIS

Abatacept
Brand(s): Orencia
DOSAGE FORM/ STRENGTH: 250 mg/15 mL intravenous injection, 125 mg/mL prefilled syringe for subcutaneous injection

For the treatment of adult patients with severe active rheumatoid arthritis who meet the following criteria:

The Patient has severe active disease as demonstrated by:

- ≥ 5 swollen joints; AND
- rheumatoid factor positive; AND/OR
- having radiographic evidence of rheumatoid arthritis

Despite the optimal* use of various disease-modifying anti-rheumatic drugs (“DMARDs”).

*For the purpose of the criteria, the optimal use of DMARDs is defined as:

- use of methotrexate (dosed at 20 mg per week) for at least 3 months; AND
- use of leflunomide (dosed at 20 mg per day) for at least 3 months; AND
- an adequate trial (3 months) of at least one combination of DMARDs;
  OR
- use of methotrexate (dosed at 20 mg per week) for at least 3 months; AND
- leflunomide in combination with methotrexate for at least 3 months.

Note: If the patient cannot be treated with adequate trial(s) of methotrexate and/ or leflunomide due to contraindication(s) or intolerance(s), the nature of the contraindication(s) or intolerance(s) must be provided along with details of trials of other DMARDs or clear rationale why other DMARDs cannot be considered.

For patients who have failed treatment with an anti-TNF therapy due to lack of efficacy or toxicity, prescribers should consider use of a biologic with a different mechanism of action.
Abatacept
Brand(s): Orencia
DOSAGE FORM/ STRENGTH: 250 mg/15 mL intravenous injection, 125 mg/mL pre-filled syringe for subcutaneous injection

Approved Dosing:

IV use: The initial dose is administered at 0, 2, and 4 weeks then every 4 weeks thereafter. Note that funding for higher doses will not be considered.

<table>
<thead>
<tr>
<th>Body weight of patient</th>
<th>Dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>&lt; 60 kg</td>
<td>500 mg</td>
</tr>
<tr>
<td>60-100 kg</td>
<td>750 mg</td>
</tr>
<tr>
<td>&gt;100 kg</td>
<td>1 gram</td>
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</tbody>
</table>

SC use: 125 mg SC weekly. Note that an IV loading dose of 750 mg may be given prior to initiating the weekly SC dosing. (Please refer to the Orencia product monograph for further details.)

Duration of Approval: First Renewal – 1 Year, Subsequent Renewals – 5 Years

Renewals will be considered in patients with objective evidence of at least a twenty percent (20%) reduction in swollen joint count and a minimum of improvement in two (2) swollen joints over the previous year.

For renewals beyond the second year, objective evidence of the preservation of treatment effect must be provided by the requesting physician.
**Adalimumab – see Formulary for funded biosimilars**
Brand(s): Humira and formulary listed biosimilars
DOSAGE FORM/ STRENGTH: 40 mg/0.8mL prefilled syringe, 40 mg/0.8mL and 20 mg/0.2 mL prefilled pens for subcutaneous injection

**Anakinra**
Brand(s): Kineret
DOSAGE FORM/ STRENGTH: 100 mg /0.67 mL subcutaneous injection

**Certolizumab pegol**
Brand(s): Cimzia
DOSAGE FORM/ STRENGTH: 200 mg/mL prefilled syringe and autoinjector

**Etanercept – see Formulary for funded biosimilars**
Brand(s): Enbrel
DOSAGE FORM/ STRENGTH: 25 mg and/or 50 mg prefilled syringe or pens for subcutaneous injection per formulary listed options

**Golimumab**
Brand(s): Simponi
DOSAGE FORM/ STRENGTH: 50 mg/0.5 mL prefilled syringe and autoinjector

**Infliximab – see Formulary for funded biosimilars**
Brand(s): Remicade
DOSAGE FORM/ STRENGTH: 100 mg/vial intravenous infusion

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**For the treatment of rheumatoid arthritis** in patients who have:
- Severe active disease (≥ 5 swollen joints and rheumatoid factor positive and/or, anti-CCP positive, and/or radiographic evidence of rheumatoid arthritis) despite the optimal use of various formulary disease-modifying anti-rheumatic drugs (DMARDs)*.  

*Optimal use of DMARDs include:
- Methotrexate (20 mg/week) for at least 3 months and leflunomide (20 mg/day) for at least 3 months in addition to an adequate trial (3 months) of at least one combination of DMARDs; or
- Methotrexate (20 mg/week) for at least 3 months and leflunomide in combination with methotrexate for at least 3 months.  
- If the patient could not receive adequate trial(s) of methotrexate and/or leflunomide due to contraindication(s) or intolerance(s), the nature of contraindication(s) or intolerance(s) must be provided along with details of trials of other DMARDs or clear rationale why other DMARDs cannot be considered.

OR
• Methotrexate (20mg/week), sulfasalazine (2 GM/day) and hydroxychloroquine (400mg/day)* for at least 3 months. If the patient could not receive an adequate trial of methotrexate, sulfasalazine and hydroxychloroquine due to intolerance, then the above DMARD trial criteria must be met.

Hydroxychloroquine is based by weight up to 400 mg per day

**Duration of Approval:** 1 Year

**Renewal** will be considered for patients with objective evidence of at least a 20% reduction in swollen joint count and a minimum of improvement in 2 swollen joints over the previous year. For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided.

The planned dosing regimen for the requested biologic should be provided. The recommended doses for the treatment of rheumatoid arthritis are as follows:

- Adalimumab 40mg every two weeks
- Anakinra 100mg per day
- Certolizumab pegol 400mg at 0, 2 and 4 weeks followed by maintenance therapy of 200 mg every 2 weeks. For maintenance dosing, 400mg every 4 weeks may be considered
- Etanercept 25mg twice weekly or 50mg once weekly
- Golimumab 50mg once a month
- Infliximab 3mg/kg/dose at 0, 2 and 6 weeks followed by maintenance therapy of 3mg/kg/dose every 8 weeks up to a maximum of six maintenance doses per year

*(Note that effective December 22, 2016, Tofacitinib (Xeljanz) 5 mg is available on the ODB Formulary in patients meeting the Limited Use criteria)*

**Duration of Approval:** First Renewal – 1 Year, Subsequent Renewals – 5 Years
First course of Rituximab for the treatment of rheumatoid arthritis in adult patients with:

- Severe active disease (≥ 5 swollen joints and rheumatoid factor positive and/or radiographic evidence of rheumatoid arthritis); AND
- Failure to respond to optimal use of DMARDs or documented intolerance or contraindications to DMARDs (per current EAP reimbursement criteria for anti-TNF agents); AND
- Failure to respond to, or the patient has intolerance or contraindications to, an adequate trial of at least ONE anti-TNF agent (e.g., adalimumab, etanercept, infliximab, golimumab, certolizumab pegol)

Initial approval: One year: One course of treatment is 1000 mg followed two weeks later by the second 1000mg dose. Two courses will be approved each year (courses should be at least 6 months apart with second course being given only AFTER loss of effect as noted in the re-treatment guidelines below). Second course is not approved for “maintenance” therapy.

Renewal criteria: A joint count at 3-4 months indicating at least a 20% reduction in swollen joint count and a minimum of improvement in 2 swollen joints, should be recorded to indicate a response, and then re-treatment can be given after an interval of at least 6 months AND after a loss of effect. Details of all courses given and the subsequent response should be provided in the renewal request.

Renewal approval: 1 year (2 courses). One course of treatment is 1000 mg followed two weeks later by the second 1000mg dose. Repeated courses are not approved for maintenance therapy.

Note: Rituximab should not be used concomitantly with other anti-TNF agents.

More information describing one of the Committee to Evaluate Drugs’ review of rituximab can be found on the Ministry website.
For the treatment of rheumatoid arthritis in adult patients meeting the following criteria:

a) Sarilumab is being used as monotherapy or in combination with methotrexate or other non-biologic disease-modifying antirheumatic drugs (DMARDs); AND
b) Patient is 18 years of age or older; AND
c) Has severe active disease (≥ 5 swollen joints and rheumatoid factor positive and/or anti-CCP positive and/or radiographic evidence of rheumatoid arthritis) despite the optimal use of various formulary disease-modifying anti-rheumatic drugs (DMARDs); AND
d) Has one of the following:
   i) fails to respond to Optimal use\(^1\) of DMARDs (e.g. hydroxychloroquine, methotrexate, sulfasalazine, leflunomide, cyclosporine, azathioprine, penicillamine, chloroquine and gold compounds).

\(^1\)Optimal use of DMARDs is defined as one of the below:.

a) methotrexate (20 mg/week) for at least 3 months and leflunomide (20 mg/day) for at least 3 months, in addition to an adequate trial (3 months) of at least one combination of DMARDs;
b) methotrexate (20 mg/week) for at least 3 months and leflunomide in combination with methotrexate for at least 3 months; or
c) methotrexate (20 mg/week), sulfasalazine (2 G/day) and hydroxychloroquine (based on weight and up to 400 mg/day) for at least 3 months.

ii) has a documented intolerance or contraindication to DMARDs in which case the nature of the contraindication(s) or intolerance(s) must be provided with the request, along with details of trials of other DMARDs or clear rationale as to why other DMARDs cannot be considered

Approval duration of Initials: 1 year

Objective evidence of at least a 20% reduction in swollen joint count and a minimum of improvement in 2 swollen joints over the previous year.

Approval duration of first renewal: 1 year

For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided.

Subsequent Renewal Criteria: Approval duration 5 years
Recommended Dose:

The recommended dose of KEVZARA is 200 mg once every 2 weeks given as a subcutaneous injection.

A reduced dose of 150 mg once every two weeks is recommended for patients with neutropenia, thrombocytopenia, or with elevated liver enzymes.

**Tocilizumab**

Brand(s): Actemra

DOSAGE FORM/ STRENGTH: 80 mg/4 mL Vial, 200 mg/10 mL Vial, 400 mg/20 mL Vial, 162mg/0.9mL Inj (Prefilled syringe), 162mg/0.9mL Auto Injector

For the treatment of rheumatoid arthritis in adult patients with:

- Severe active disease (≥ 5 swollen joints and rheumatoid factor positive and/or anti-CCP positive and/or has radiographic evidence of rheumatoid arthritis); **AND**

- Failure to respond to optimal use\(^1\) of DMARDs or with documented intolerance to DMARDs (per current EAP reimbursement criteria for anti-TNF agents).

Optimal use of DMARDs (hydroxychloroquine, methotrexate, sulfasalazine, leflunomide, cyclosporine, azathioprine, penicillamine, chloroquine and gold compounds) defined as:

a) Methotrexate (20 mg/week) for at least 3 months **AND**

b) Leflunomide (20 mg/day) for at least 3 months, in addition to an adequate trial (3 months) of at least one combination of DMARDs; **OR**

c) Methotrexate (20 mg/week) for at least 3 months **AND** leflunomide in combination with methotrexate for at least 3 months; **OR**

\(^1\)Note: If the patient could not receive adequate trial(s) of methotrexate and/or leflunomide due to contraindication(s) or intolerance(s), the nature of the contraindication(s) or intolerance(s) must be provided along with details of trials of other DMARDs or clear rationale as to why other DMARDs cannot be considered.

d) Methotrexate (20 mg/week), sulfasalazine (2 G/day) and hydroxychloroquine (400 mg/day)\(^2\) for at least 3 months. If the patient could not receive an adequate trial of methotrexate, sulfasalazine and hydroxychloroquine due to intolerance, then the above DMARD trial criteria must be met.

\(^2\)Hydroxychloroquine is based by weight up to 400 mg per day

The requesting physician is required to provide the planned dosing regimen on the request.
**Tocilizumab**

**Brand(s):** Actemra

**DOSAGE FORM/ STRENGTH:** 80 mg / 4 mL Vial, 200 mg /10 mL Vial, 400 mg/ 20 mL Vial, 162 mg/0.9 mL solution for injection, 162 mg/0.9 mL Autoinjector

The following are the recommended doses for tocilizumab (Actemra) IV and SC for rheumatoid arthritis:

**IV recommended dose:**

Approval for 4mg/kg/dose once every 4 weeks followed by an increase to 8mg/kg/dose based on clinical response; even for individuals whose body weight is more than 100kg, doses exceeding 800mg per infusion are not recommended

**SC recommended dose:**

For patients < 100 kg weight, starting dose of 162 mg every other week, followed by an increase to every week based on clinical response. For patients at or above 100 kg weight, 162 mg every week.

**Duration of Approval:** 1 Year

**Renewal** will be considered for patients with objective evidence of at least a 20% reduction in swollen joint count and a minimum of improvement in 2 joints over the previous year.

For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided.

Duration of Approval of first Renewal – 1 Year

Duration of Second and Subsequent Renewals – 5 Years
Substance Dependence

Acamprosate Calcium
Brand(s): Campral
DOSAGE FORM/ STRENGTH: 333 mg tablet

Note: Effective June 29, 2018, Acamprosate is made available as a Limited Use drug on the Ontario Drug Benefit Formulary.

Methadone Compounded Solution
Brand(s):
DOSAGE FORM/ STRENGTH:

Effective September 1, 2014

Reimbursement of Compounded Methadone solution for the treatment of opioid dependence will be considered for patients who meet the following criteria;

Patient has demonstrated that they have experienced a true allergy to both commercially available Methadose formulations (i.e., Methadose 10 mg/mL oral cherry flavoured concentrate AND Methadose 10 mg/mL dye-free, sugar-free, unflavoured oral concentrate).

The request must be accompanied by a completed Health Canada adverse drug reaction form (Canada Vigilance Adverse Reaction Reporting Form) and include a detailed description of the allergic reaction to each Methadose product, a description of the circumstances in which the reactions occurred, and demonstration that the allergy is unlikely to be related to any diluent in which Methadose was mixed, but rather, that it was caused by the excipients within the Methadose formulation.

Naltrexone
Brand(s): Revia
DOSAGE FORM/ STRENGTH: 50 mg tablet

Note: Effective June 29, 2018, Naltrexone is made available as a Limited Use drug on the Ontario Drug Benefit Formulary.
SYSTEMIC JUVENILE IDIOPATHIC ARTHRITIS

Anakinra
Brand(s): Kineret
DOSAGE FORM/ STRENGTH: 100mg/0.67mL pre-filled syringe

For the treatment of systemic juvenile idiopathic arthritis in patients who meet the following criteria;

a) Patient must have a diagnosis of sJIA with fever ( >38 degrees Celsius) for at least 2 weeks AND at least ONE of the following:
   - rash of systemic JIA
   - serositis (e.g. pericarditis, pleuritis, or peritonitis)
   - lymphadenopathy (e.g. cervical, axillary, inguinal)
   - hepatomegaly
   - splenomegaly

b) The physician making the request has ruled out other potential etiologies (e.g. malignancies, serious clinical infections, and other inflammatory or connective tissue diseases); AND

c) Age of disease onset is younger than 16 years of age. (Note: the physician must specify age of disease onset in the request); AND

d) Systemic corticosteroids cannot be used for at least ONE of the following reasons (please specify name and current dose of corticosteroid, if applicable):
   - The patient is unresponsive and/or refractory to systemic corticosteroids; OR
   - The patient has experienced a systemic reaction (e.g. fever, rash of sJIA, serositis, lymphadenopathy, hepatomegaly or splenomegaly) while on tapering doses of systemic corticosteroids (i.e. the patient is corticosteroid dependent); OR
   - The patient has experienced an adverse drug reaction to a systemic corticosteroid; OR
   - The use of systemic corticosteroids is contraindicated in this patient.
Anakinra
Brand(s): Kineret
DOSAGE FORM/ STRENGTH: 100mg/0.67Ml pre-filled syringe

Note: The following requests will undergo external review on a case-by-case basis:

- Patients with Macrophage Activation Syndrome
- Patients who meet initial sJIA criteria and are currently 16 years of age or older
- Patients who meet initial sJIA criteria and are requesting higher dosing regimens
  (Please provide rationale for the higher dosing regimen with your request)

Dosing: 1-2 mg/kg subcutaneously once daily.

Duration of Approval: 1 Year

Renewal will be considered for patients demonstrating at least a 50% reduction in corticosteroid dose (unless contraindicated, not tolerated, unresponsive or refractory at the time of initial request) and no evidence of active systemic disease. For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided.

The following renewal requests will undergo external review:

- Evidence of active systemic disease
- Requests for higher dosing regimens (Please provide rationale for the higher dosing regimen with your request)
- Patient is currently 16 years of age or older

Duration of Approval: 1 Year
For the treatment of systemic juvenile idiopathic arthritis in patients who meet the following criteria;

- Patient must have a diagnosis of sJIA with fever (>38 degrees Celsius) for at least 2 weeks AND at least ONE of the following:
  - rash of systemic JIA
  - serositis (e.g. pericarditis, pleuritis, or peritonitis)
  - lymphadenopathy (e.g. cervical, axillary, inguinal)
  - hepatomegaly
  - splenomegaly

- The physician has ruled out other potential etiologies (e.g. malignancies, serious clinical infections, and other inflammatory or connective tissue diseases); AND
- Age of disease onset is younger than 16 years of age. (Note: the physician must specify age of disease onset in the request); AND
- Systemic corticosteroids cannot be used for at least ONE of the following reasons (please specify name and current dose of corticosteroid, if applicable):
  - The patient is unresponsive and/or refractory to systemic corticosteroids; OR
  - The patient has experienced a systemic reaction (e.g. fever, rash of sJIA, serositis, lymphadenopathy, hepatomegaly or splenomegaly) while on tapering doses of systemic cortico-steroids (i.e. the patient is corticosteroid dependent); OR
  - The patient has experienced an adverse drug reaction to a systemic corticosteroid; OR
  - The use of systemic corticosteroids is contraindicated in this patient.

Note: The following requests will undergo external review on a case-by-case basis:

- Patients with Macrophage Activation Syndrome
- Patients who meet initial sJIA criteria and are currently 16 years of age or older
- Patients who meet initial sJIA criteria and are requesting higher dosing regimens (Please provide rationale for the higher dosing regimen with your request)
Tocilizumab
Brand(s): Actemra
DOSAGE FORM/ STRENGTH: 80 mg / 4 mL, 200 mg / 10 mL, 400 mg / 20 mL, 162 mg sc inj (Prefilled Syringe), 162 mg Autoinjector

Dosing: For those less than 30 kg, 12 mg/kg IV every 2 weeks
For those greater than or the same as 30 kg, 8 mg/kg IV every 2 weeks

Note: Recommended maximum adult dose is 800 mg.

SC dosing regimen:

a) 162 mg once every 2 weeks if the Patient weighs less than 30 kg
b) 162 mg once every week if the Patient weighs 30 kg or more

Duration of Approval: 1 Year

Renewal will be considered for patients demonstrating at least a 50% reduction in corticosteroid dose (unless contraindicated, not tolerated, unresponsive or refractory at the time of initial request) and no evidence of active systemic disease. For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided.

The following renewal requests will undergo external review:
- Evidence of active systemic disease
- Requests for higher dosing regimens (Please provide rationale for the higher dosing regimen with your request)

Patient is currently 16 years of age or older

Duration of Approval: 1 Year
JUVENILE SPONDYLOARTHRITIS OR ENTHESITIS-RELATED ARTHRITIS

Adalimumab – see Formulary for funded biosimilars
Brand(s): Humira and formulary listed biosimilars
DOSAGE FORM/STRENGTH: 40mg/0.8mL prefilled syringe, 40mg/0.8mL and 20mg/0.2mL prefilled pens for subcutaneous injection

Etanercept – see Formulary for funded biosimilars
Brand(s): Enbrel and formulary listed biosimilars
DOSAGE FORM/STRENGTH: 25mg/vial, 50 mg prefilled syringe for subcutaneous injection

Infliximab – see Formulary for funded biosimilars
Brand(s): Remicade and formulary listed biosimilars
DOSAGE FORM/STRENGTH: 100 mg/vial

Updated: March 29, 2021

For the treatment of juvenile spondyloarthritis (JSpA) or enthesitis-related arthritis (ERA) in patients who meet the following criteria for either axial or peripheral disease:

Axial Disease

- Age of disease onset ≤ 16 years; AND
- Low back pain and stiffness for > 3 months that improve with exercise and not relieved by rest; AND
- Failure to respond to or documented intolerance to adequate trials of 2 nonsteroidal anti-inflammatory drugs (NSAIDs) for at least 4 weeks each; AND
- BASDAI score of ≥ 4 after at least 4 weeks of standard NSAID therapy; AND
- Imaging evidence of severe active disease by X-ray, CT scan or MRI

The details of imaging reports for severe active disease must provide the following:
  - X-ray or CT scan report stating the presence of “SI joint fusion” or “SI joint erosion” OR
  - MRI report stating the presence of “inflammation” or “edema” or “erosion” of the SI joint.

Actual imaging reports must be submitted with the request. If the imaging reports do not specify the above findings, the request will be reviewed by external medical experts. The imaging interpretation report from the radiologist or rheumatologist may be submitted along with radiographic report.
**Adalimumab** – see Formulary for funded biosimilars  
Brand(s): Humira and formulary listed biosimilars  
DOSAGE FORM/ STRENGTH: 40 mg/0.8 mL prefilled syringe, 40mg/0.8mL and 20 mg/0.2 mL prefilled pen for subcutaneous injection

**Etanercept** – see Formulary for funded biosimilars  
Brand(s): Enbrel and formulary listed biosimilars  
DOSAGE FORM/ STRENGTH: 25mg/vial, 50 mg prefilled syringe for subcutaneous injection

**Infliximab** – see Formulary for funded biosimilars  
Brand(s): Remicade and formulary listed biosimilars  
DOSAGE FORM/ STRENGTH: 100 mg/vial

**Peripheral Disease**

- Age of disease onset ≤ 16 years; AND  
- ≥ 5 active sites of inflammation attained by a combination of swollen/active joints and/or enthesitis sites (tenderness or swelling at enthesal insertion)  
- Failure or intolerance to at least one DMARD (sulfasalazine 50 mg/kg/day-maximum 2 grams, or methotrexate 15 mg/m²/week-maximum 25 mg per week) for at least 3 months.

**Renewals** will be considered for patients with objective evidence of at least a 50% reduction in BASDAI score or ≥ 2 absolute point reduction in BASDAI score.

For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided.

**Renewal** will be considered for patients with objective evidence of at least a 20% reduction in active sites over the previous year. There should also be an improvement in number of enthesitis sites.

For renewals beyond the second year, objective evidence of preservation of treatment effect must be provided. Requests that do not meet these criteria will undergo external review.

The planned dosing regimen for the requested biologic should be provided.

The recommended dose for the treatment of JSpA/ERA is as follows:

i) Etanercept 0.4mg/kg (max 25 mg) twice weekly or 0.8mg/kg (max 50 mg) once weekly
ii) Infliximab: 5mg/kg/dose at 0, 2 and 6 weeks followed by maintenance therapy of up to 5mg/kg/dose every 6-8 weeks

iii) Adalimumab:
    If Less than 30 kg: 20 mg SC q 2 weeks
    If Greater than or equal to 30 kg: 40 mg SC every 2 weeks

Requests for higher doses will be considered on a case-by-case basis.

Duration of Approval of Initials and Renewals: 1 Year

**SPASTICITY TREATMENTS**

**Tizanidine**
Brand(s): Zanaflex
DOSAGE FORM/ STRENGTH: 4 mg tablet

For the treatment of spasticity in patients who have failed and/or cannot tolerate at least two of the following available alternatives: baclofen, diazepam and dantrolene.

- Submission must describe the intolerance experienced.

Duration of Approval: Lifetime

**URINARY ANTISPASMODICS**

**Oxybutynin Transdermal System**
Brand(s): Oxytrol
DOSAGE FORM/ STRENGTH: 36 mg transdermal patch (3.9 mg/day system)

The treatment of urinary frequency, urgency or urge incontinence in patients who are unable to take oral treatments (e.g. inability to swallow or who are unable to absorb (e.g. short gut syndrome).

Adverse effects to oral therapy (e.g. dizziness) are not acceptable.

Duration of Approval: 5 Years
EXCEPTIONAL ACCESS PROGRAM

TELEPHONE REQUEST SERVICE (TRS)

REIMBURSEMENT CRITERIA FOR SELECTED TRS DRUGS
INTRODUCTION

The Ontario Public Drug Programs has developed these reimbursement criteria to provide physicians with information about selected drug products that may be considered for funding through the Exceptional Access Program’s Telephone Request Service (TRS). The TRS offers prescribers (i.e. physicians and nurse practitioners) another way to submit EAP requests for a group of selected drugs. This document provides a list of the drugs and their funding criteria that are considered through the TRS. In most cases, the request will be assessed during the call and a funding decision provided to the caller by the end of the call. In general, approvals will be processed within one business day turnaround.

Prescribers (or their delegates) are encouraged to review the reimbursement criteria for the drug being requested before calling the service to ensure that all of the necessary information is available during the call. Callers who wish to submit a request for drug products and indications not currently available through TRS will be asked to fax the request to EAP. If your request is approved, the prescriber will receive a response letter notifying him/her of the funding decision.

The EAP response letter will list the specific drug, drug identification number (DIN) or product identification number (PIN), strength and dosage form that is considered for funding. Prescribers and pharmacists are responsible to ensure that funded products are provided to avoid unnecessary out-of-pocket costs to the patient. Note that not all generic brands are funded or interchangeable (on-formulary or off-formulary). You can also refer to the formulary for a list of the most updated interchangeable drugs products.

The Ministry reserves the right to change the list of drug products at its sole discretion. If you have any questions or concerns regarding the TRS, please contact us at:

Exceptional Access Program – Telephone Request Service 3rd Floor, 5700 Yonge St. North York, ON M2M 4K5 Phone: 1-866-811-9893 or 416-327-8109 Fax: 1-866-811-9908 or 416-327-7526

E-mail: EAPFeedbackLine@ontario.ca
**Anti-Infectives**

**Cefazolin**  
**Brand(s):** Many Generic Brands  
**DOSAGE FORM/ STRENGTH:** 1 g/vial Injection  

For treatment of infections susceptible to cefazolin.  

**Standard Approval Duration:** As requested up to 5 years

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**Daptomycin**  
**Brand(s):** Cubicin RF, daptomycin generics (refer to formulary for funded drugs)  
**DOSAGE FORM/ STRENGTH:** 500 mg/10mL injection  

For the treatment of patients with one or more of the following condition(s):  

1. Osteomyelitis caused by methicillin-resistant staphylococcus aureus (MRSA)  
2. Device-related osteoarticular or prosthetic joint infections caused by methicillin-resistant staphylococcus aureus (MRSA);  
3. Diabetic foot infections caused by methicillin-resistant staphylococcus aureus (MRSA); AND/OR  
4. Staphylococcus aureus bloodstream (SAB) infection including right-sided  
   Staphylococcus aureus infective endocarditis (SARIE) infection caused by methicillin-resistant Staphylococcus aureus (MRSA)

Additionally, the patient must have failed to adequately respond to, be intolerant* to, or have a contraindication to vancomycin.

*Requests involving red-man-syndrome with vancomycin must provide details of the intolerance including the rate of infusion and the use of antihistamines and other histamine blockers prior to therapy.

**Standard Approval Duration:** Up to maximum of 56 days

**Exclusion Criteria:** Daptomycin is not funded for patients with:  
   a) MRSA-related pneumonia;  
   b) ii) skin/skin structure infections other than diabetic foot infections caused by MRSA.
Daptomycin
Brand(s): Cubicin RF, generic daptomycin (refer to the formulary for funded drugs)

DOSAGE FORM/STRENGTH: 500 mg/10 mL powder for injection

For the treatment of invasive infections\(^1\) caused by vancomycin-resistant enterococcus (VRE) in patients who meet the following criteria:

1. VRE infection is confirmed by blood or tissue culture and sensitivity report
2. Patient is unable to use linezolid as a result of at least ONE of the following reasons:
   i) has developed resistance to linezolid as confirmed by the microbiology sensitivity report
   ii) has experienced intolerance\(^2\) to linezolid (for example: severe gastric symptoms, myelosuppression, peripheral neuropathy requiring medical intervention, lactic acidosis)
   iii) has a contraindication\(^2\) to linezolid

OR

Prescribed by an infectious disease expert for a patient who is able to use linezolid but where the bacteriostatic effect of linezolid may not be deemed to be clinically optimal due to other patient factors or comorbidities (e.g., immunocompromised, neutropenic).

\(^1\)Not approved for colonization (e.g., nares, skin, stool)

\(^2\)Intolerances and contraindications to linezolid must be fully described in the EAP application.

Recommended dose: 8 to 12 mg/kg daily for VRE with adjustments based on renal function

Duration of approval:

*Note that the below are examples of durations for reimbursement of some common VRE infections. This is not an all-inclusive list.*
Urinary tract infections: up to 10 days
Bacteremia: up to 14 days
Endocarditis: up to 6 weeks
Osteomyelitis: up to 8 weeks

Requests for longer durations of funding will be considered case-by-case through external review and must faxed to the EAP and be accompanied by a recent microbiology sensitivity report to confirm sensitivity to daptomycin.

Requests for daptomycin for other types of infections may be faxed to the EAP for case-by-case consideration.
Fidaxomicin
Brand(s): Dificid
DOSAGE FORM/ STRENGTH: 200 mg tablet

For the treatment of Clostridium difficile infection (CDI) in patients who meet the EAP criteria for vancomycin use, but 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 failure* 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 intolerance** to oral vancomycin treatment that resulted in the discontinuation of vancomycin therapy.

*Treatment failure is defined as 7 days of vancomycin therapy without acceptable clinical improvement.

**Details of severe adverse reaction or intolerance must be provided and should be clinically related to oral administration of vancomycin.

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: Fecal biotherapy (“stool transplantation”), if available, should be encouraged for this patient population.

Approved dose and duration: 200 mg twice a day for 10 days

Gentamycin
Brand(s): Many Generic Brands
DOSAGE FORM/ STRENGTH: 40 mg/mL injection

For treatment of infections susceptible to gentamycin.

Standard Approval Duration: As requested up to 5 years
Isavuconazole
Brand(s): Cresemba
DOSAGE FORM/ STRENGTH: 100 mg capsule, 200 mg Injection Solution

NOTE: Prescribers must submit the laboratory results to confirm the patient's infection and include drugs and drug regimens that have been used for the patient's condition, including the response to prior therapies and other relevant clinical information. Please also confirm that the patient does not meet the exclusion criteria.

For the treatment of invasive aspergillosis in patients meeting the following criteria:

- 18 years of age and older; AND
- Patient has failed, experienced intolerance to, or has contraindications to voriconazole; AND
- Isavuconazole is prescribed by or in consultation with an infectious disease specialist.
  (Include consult note with the request)

Exclusion criteria:

Patients with familial short QT syndrome.

Approval Duration: 3 months

Renewals will be considered on a case-by-case basis.

For the treatment of invasive mucormycosis in patients meeting the following criteria:

- 18 years of age and older; AND
- Isavuconazole is prescribed by or in consultation with an infectious disease specialist.
  (Include consult note with the request)

Exclusion criteria:

Patients with familial short QT syndrome.

Approval Duration: 3 months

Renewals will be considered on a case-by-case basis

Recommended dose:

200mg administered intravenously or orally every 8 hours for 6 doses followed by a maintenance dose of 200mg daily starting 12 to 24 hours after the last loading dose.

Oral therapy should be considered as a preferred option when clinically appropriate. A loading dose is not required when switching from intravenous to oral treatment or vice versa.
1. For the prophylaxis of Aspergillus and Candida infections in patients who have recently (within the past 3 months) undergone an allogeneic bone marrow transplant.

2. For the prophylaxis of invasive fungal infections in patients who have previously (3 months or longer) undergone an allogeneic stem cell transplant and are experiencing moderate to severe graft-versus-host-disease (GVHD) will be considered on a case-by-case basis.

   **Renewals** will be considered on a case-by-case basis for patients who continue to experience ongoing symptoms of moderate to severe GVHD. Please provide information regarding infections that were experienced while on therapy (as applicable) including the names of medications and treatments being used to manage GVHD.

3. For the treatment of invasive aspergillosis* in patients who are refractory or intolerant to voriconazole OR who have documented contraindication to voriconazole.

   *Invasive aspergillosis should be confirmed by fungal culture.

   Note: Requests without a positive fungal culture must be accompanied by a consultation note from an infectious disease expert with details of how the diagnosis was made and will be considered on a case-by-case basis.

   **Renewals** will be considered on a case-by-case basis.

4. For the treatment of mucormycosis** in patients who have failed, have a contraindication to, or experienced intolerance to amphotericin B; OR

   For the step-down treatment of mucormycosis** in patients who have been initially treated with amphotericin B but cannot tolerate long-term therapy with this agent.

   **Mucormycosis infection must be confirmed by fungal culture.

Note: Requests without a positive fungal culture but where the diagnosis of mucormycosis is documented by an infectious diseases consult and other tools (e.g., radiology reports, histopathology, etc.) will be considered on a case-by-case basis.

**Renewals** will be considered for patients who are responding to therapy but who have not experienced clinical resolution of their condition. Note that requests for renewal must be accompanied by supporting clinical information (Infectious disease consultation/radiology report).
# Chronic Renal Failure Drugs

## Calcium Carbonate
*Brand(s): Tums*

**DOSAGE FORM/ STRENGTH:** 500 mg, 750 mg, 1000 mg

For patients with hypoparathyroid disease or chronic renal failure.

*NOTE: Calcium supplements for patients who do not have hypoparathyroid disease or chronic renal failure are not eligible for funding consideration by the ODB program, which includes EAP.*

Renewals will be considered where patient is stable.

**Standard Approval Duration:** 5 years for initials and renewals

## Lanthanum
*Brand(s): Fosrenol and Generics*

**DOSAGE FORM/ STRENGTH:** 250 mg, 500 mg, 750 mg, 1000 mg Chewable tablet

## Sevelamer Hydrochloride
*Brand(s): Renagel*

**DOSAGE FORM/ STRENGTH:** 800 mg tablet

## Sevelamer Carbonate
*Brand(s): Accel-Sevelamer*

**DOSAGE FORM/ STRENGTH:** 800 mg tablet

## Sevelamer Carbonate
*Brand(s): Renvela*

**DOSAGE FORM/ STRENGTH:** 800 mg tablet, 0.8 g powder sachet, 2.4 g powder sachet

*Updated: June 7, 2021*

i) For the treatment of hyperphosphatemia associated with end-stage renal disease (ESRD) where patients are on dialysis and have a sustained serum phosphate > 1.8 mmol/L AND adjusted serum calcium > 2.65 mmol/L; OR

ii) For dialysis patients experiencing hyperphosphatemia (sustained serum phosphate levels >1.8 mmol/L) who have calciphylaxis and/or evidence of coronary artery calcification.
Notes:
1. Calcium and phosphate levels provided to demonstrate sustained elevations should be at least 4 weeks apart.

2. Patients who demonstrate reduction of phosphate levels while on a calcium binder but who experience adjusted serum calcium > 2.65 mmol/L must provide supporting laboratory levels before treatment and while on treatment and should include the dose(s) of calcium-based binder used.

**Exclusion Criteria:**
Sevelamer will not be reimbursed in the following cases:
Use in combination therapy with another funded non-calcium-based phosphate binder (e.g. lanthanum carbonate hydrate, sucroferric oxyhydroxide, other sevelamer formulations)

**Duration of approval:** lifetime

On a case-by-case basis, requests may be considered for sevelamer under the EAP for a patient with serum phosphate less than 1.8mmol/L and calcium values less than 2.65 mmol/L.

**Sucroferric Oxyhydroxide**
Brand(s): Velphoro
**DOSAGE FORM/ STRENGTH:** 500 mg iron per chewable tab

For the treatment of hyperphosphatemia associated with end-stage renal disease (ESRD) where patients are on dialysis and have a sustained serum phosphate > 1.8 mmol/L AND serum calcium > 2.65 mmol/L; OR

For dialysis patients experiencing hyperphosphatemia (sustained serum phosphate levels >1.8 mmol/L) who have calciphylaxis and/or evidence of coronary artery calcification.

Note: Dialysis patients with hyperphosphatemia meeting the above criteria who are experiencing other types of calcification (e.g. aortic) in the absence of coronary artery calcification may be considered case-by-case by faxing your request to the EAP.

Calcium and phosphate levels provided to demonstrate sustained elevations should be at least 4 weeks apart.

**Exclusion criteria:**
Patients with haemochromatosis or any other iron accumulation disorders.

**Duration of Approval:** Lifetime
Vitamin B Complex with Vitamin C
Brand(s): Replavite plus Generics
DOSAGE FORM/ STRENGTH:

For patients receiving hemodialysis or peritoneal dialysis.

Standard Approval Duration: 5 years
Anticoagulants

NOTE: Low Molecular Weight Heparins (LMWHs) are currently listed on the ODB Formulary as Limited Use (LU) benefits for the treatment of deep venous thrombosis (DVT) and pulmonary embolism (PE) in certain patient groups. Please consult the Formulary for further details.

Dalteparin
Brand(s): Fragmin
DOSAGE FORM/ STRENGTH: Check e-formulary for funded strengths

1. For peri-operative bridging for patients who require long-term warfarin therapy and must temporarily discontinue it before and after surgery, and who are at moderate- to high-risk for an embolic event while off warfarin.,

   Standard Approval Duration: As requested up to a maximum of 10 days before the date of surgery plus up to 7 days after the date of hospital discharge

2. For post-operative prophylaxis of DVT for patients who had hip or knee surgery and cannot use warfarin. 10

   Standard Approval Duration: As requested up to a maximum of 30 days starting on the day of surgery.

3. For the post-operative prophylaxis of venous thromboembolism following abdominal or pelvic surgery for cancer in patients who do not have a history of or risk factors for heparin-induced thrombocytopenia.

   Standard Approval Duration: Maximum of 30 days.

4. For extended treatment of symptomatic acute venous thromboembolism (VTE) in patients with cancer, who cannot use warfarin.

   Standard Approval Duration: As requested up to 6 months
1. For **peri-operative bridging** for patients who require long-term warfarin therapy and must temporarily discontinue it before and after surgery, and who are at moderate- to high-risk for an embolic event while off warfarin.,

   **Standard Approval Duration**: As requested up to a maximum of **10 days** before the date of surgery **plus up to 7 days** after the date of hospital discharge

2. For **post-operative prophylaxis of DVT** for patients who had hip or knee surgery and cannot use warfarin. 10

   **Standard Approval Duration**: As requested up to a maximum of **30 days** starting on the day of surgery.

3. For the **post-operative prophylaxis of venous thromboembolism following abdominal or pelvic surgery for cancer** in patients who do not have a history of or risk factors for heparin-induced thrombocytopenia.

   **Standard Approval Duration**: Maximum of 30 days.
Oral Hypoglycemic Agents

Note: Prescribers do not need to make an EAP request for patients currently receiving pioglitazone or rosiglitazone through ODB. Physicians will be required to make an application for coverage for any patient new to ODB that is being started on either of these drugs or any ODB recipient who is new to using these drugs.

Requests for ongoing treatment with pioglitazone or rosiglitazone for patients who were previously covered by other means may be considered according to renewal criteria.

Funding under the EAP for pioglitazone or rosiglitazone will not be provided in the following clinical settings:

- Patients with type 1 diabetes
- Monotherapy, even if patient is intolerant or has contraindications to both metformin and sulfonylureas
- Combination use with either nitrates or insulin
- Patients with any stage of heart failure (NYHA Class I, II, III, IV)
- Patients at high risk for bone fracture (post-menopausal women with previously confirmed osteoporosis or osteopenia)
- Patients with recent history (in the past 3 months) of ischemic cardiovascular event (myocardial infarction, unstable angina)
- Patients with active bladder cancer, a history of bladder cancer or uninvestigated macroscopic haematuria

### Pioglitazone

**Brand(s):** Generics

**DOSAGE FORM/ STRENGTH:** 15 mg, 30 mg, 45 mg tablet

For dual combination therapy of type 2 diabetes, in patients with:

a) Inadequate glycemic control (HbA1c of >7%) on maximal doses of metformin (2000 mg/day); OR

b) Inadequate glycemic control, on maximal doses of sulfonylurea (glyburide 10mg/day, gliclazide 160mg/day or gliclazide modified release (MR) 60 mg/day) or glimepiride 4 mg/day and demonstrated intolerance / contraindication to metformin

For triple combination therapy of type 2 diabetes, in patients with:

a) Inadequate glycemic control on maximal doses of metformin and a sulfonylurea AND only if:

- physician has offered insulin as alternative option first, and patient has refused or is not able to take insulin, AND both physician and patient are aware that thiazolidinediones are not indicated for use in triple therapy.

**Standard Approval Duration:** 5 years
**Pioglitazone**
**Brand(s):** Generics  
**DOSAGE FORM/ STRENGTH:** 15mg, 30 mg, 45 mg tablet

Renewals: EAP will renew pioglitazone only for patients who have achieved adequate glycemic control (HbA1c of ≤ 7% while on therapy and who have no known contraindications to pioglitazone.

**Standard Approval Duration:** 5 years

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**Rosiglitazone**
**Brand(s):** Avandia  
**DOSAGE FORM/ STRENGTH:** 2 mg, 4 mg, 8 mg tablet

For the treatment of type 2 diabetes mellitus in patients with:

- Inadequate glycemic control (HbA1c >7%) from ALL other oral antidiabetic agents* funded through one of the Ontario Drug Benefit (ODB) Programs, in monotherapy or in combination OR
- Where ALL other oral antidiabetic agents are inappropriate due to contraindications or intolerance AND
- The patient has refused or is not able to take insulin AND
- There is no known contraindication to rosiglitazone.

* Oral antidiabetics that need to be tried prior to consideration of rosiglitazone include the following agents currently reimbursed through the Ontario Public Drug Programs;
  - glyburide
  - metformin
  - gliclazide (Diamicron, Diamicron MR)
  - sitagliptin (Januvia)
  - repaglinide (GlucoNorm)
  - pioglitazone (Actos)
  - saxagliptin (Onglyza)

Note: A trial with acarbose is not a mandatory requirement.

Note: It is not necessary for patients to have tried the following oral antidiabetic agents that are currently not funded by the Ontario Public Drug Programs for the purposes of obtaining rosiglitazone:

**Standard Approval Duration:** 5 years
**Rosiglitazone**  
*Brand(s): Avandia*  
*DOSAGE FORM/ STRENGTH: 1 mg, 4 mg, 8 mg tablet*

Renewals will be considered where patients have benefited and continue to benefit from rosiglitazone treatment as demonstrated by achieving adequate glycemic control. This is shown by a recent HbA1c levels ≤7% while on treatment with rosiglitazone AND in who continue to have no known contraindication(s) to rosiglitazone.

**Standard Approval Duration:** 5 years

**Palliative Care Medications**

NOTE: Specific products used to treat ODB-eligible patients undergoing palliative care are reimbursed under the Ontario Public Drug Programs, as Limited use benefits on the ODB formulary or through the Facilitated Access process. Under this process, a select group of participating physicians and nurse practitioners are exempt from obtaining approval under EAP on a case-by-case basis. This assumes that the prescriber has met the qualifications set by their professional associations who administer the enrollment of their members. The prescriber’s license number with their regulatory body must appear on the prescription, for purposes of verification.

Palliative Care medication claims to be reimbursed by the ODB program must be prescribed in accordance with the following patient eligibility criteria: “This patient has a progressive, life-limiting illness and has chosen outpatient palliative treatment. Life expectancy of one year is applied to request durations.

In order to participate in the Facilitated Access to Palliative Care Drugs process, these prescribers must be registered with their professional association as meeting the qualifications for pCFA enrollement. For physicians this is by the Ontario Medical Association ("OMA") and must meet pre-defined criteria the OMA sets. For nurse practitioners, this may be the NPAO or the Nurse practitioners Association of Ontario (NPAO) or the RNAO, th Registered Nurse Association of Ontario. To facilitate the reimbursement process at the pharmacy, these prescribers are asked to indicate either, "Palliative" or “P.C.F.A.” on the prescription.

Prescribers who are not registered through this process must obtain approval through the Exceptional Access Program. A prescriber must provide the details of the patient’s diagnosis, current clinical status, and life expectancy.

For further information regarding the list of physicians and/or the criteria physicians require to be included on the list, please contact the Ontario Medical Association: (416) 340-2234, or via email. The following products can be reimbursed for the management of patients receiving palliative care through the Telephone Request Service. Note that many Palliative Care drugs have tranoptisioned to the ODB formulary for funding under Limited Use and do not require EAP authorization.
**Methadone**
Brand(s): Metadol  
DOSAGE FORM/ STRENGTH: 1 mg/mL oral liquid, 10 mg/mL oral liquid, 1 mg, 5 mg, 10 mg 25 mg  

If traditional narcotic analgesics fail to control pain or lead to side effects.  
**Standard Approval Duration:** 12 months  

**Morphine**  
Brand(s): Doral 1, Doral 5 (TEMPORARY BENEFIT)  
DOSAGE FORM/ STRENGTH: 1 mg/mL oral liquid, 5 mg/mL oral liquid  

For use in a palliative patient requiring oral liquid morphine. Doral oral liquid products are interim/temporary listing authorized by the Executive Officer as oral liquid morphine products on the formulary were withdrawn from the market by the manufacturer.  

**Oxycodone**  
Brand(s): Supeudol  
DOSAGE FORM/ STRENGTH: 5 mg, 10 mg, 20 mg  

For use when palliative patient cannot use combination oxycodone and acetaminophen.  
**Standard Approval Duration:** 12 months  

**Oxycodone HCl Controlled Release**  
Brand(s): OxyNEO  
DOSAGE FORM/ STRENGTH: 10 mg, 15 mg, 20 mg, 30 mg, 40 mg tablets  

For the treatment of cancer-related pain or pain in patients receiving end-of-life palliative care AND the patient has experienced intolerance or has failed an adequate trial (for example, three months) of at least one other listed long-acting opioid product.  
**Standard Approval Duration:** 12 months  

**Pamidronate**  
Brand(s): Many Generics  
DOSAGE FORM/ STRENGTH: 3 mg/mL, 6 mg/mL, 9 mg/mL  

For the treatment of tumor-induced/malignancy-related hypercalcemia in a palliative care patient.  
**Standard Approval Duration:** 12 months
High Dose Opioids

Effective January 31, 2017, meperidine 50mg tabs and the higher strengths of long-acting opioids including: morphine SR 200mg tabs; hydromorphone CR 24mg and 30 mg caps and fentanyl 75mcg/hr and 100mcg/hr patches were delisted from the ODB Formulary.

Access to the higher strengths of long-acting opioids is currently maintained for patients requiring palliative care through the ODB program’s:

1. Palliative Care Facilitated Access (PCFA) mechanism, for prescribers (physicians or nurse practitioners) who are registered PCFA prescribers with their professional associations; AND

2. Exceptional Access Program (EAP) Telephone Request Service (TRS) for physicians who are not PCFA prescribers, according to specific criteria.

   i) Use of the high-strength opioid must be for a patient considered to have a progressive life-limiting illness requiring palliative care.
   ii) The use of the high-strength opioid can be for pain or for symptom management.
   iii) Prescriber must have a consult from a PCFA-registered prescriber with OMA; CPSO of the PCFA-registered prescriber or NPAO or RNAO; license numbers must be provided.

**Standard Approval Duration:** 12 months

Renewals are considered with same criteria as above. A new consult from a PCFA-registered prescriber must be provided for each renewal.

Post-transplant Drugs

**Sirolimus**

Brand(s): Rapamune

**DOSAGE FORM/ STRENGTH:** 1 mg tablet, 1 mg/mL oral liquid

For liver transplant recipients who require regimens that mandate calcineurin inhibitor avoidance. The physician must be able to explain clearly why the patient cannot use a calcineurin inhibitor.

**NOTE:** Rapamune is currently listed on the ODB Formulary as a **Limited Use (LU)** benefit for the prophylaxis of organ rejection in patients receiving allogeneic renal transplants.
Valganciclovir
Brand(s): Valcyte
DOSAGE FORM/ STRENGTH: 450 mg tablet, 50 mg/mL oral solution

For the **treatment** of **cytomegalovirus (CMV) disease** following solid organ and/or bone marrow transplant in patients who meet the following criteria:

1. Objective evidence of active CMV infection determined by any one of the following methods:
   - CMV antigenemia assay; OR
   - CMV polymerase chain reaction (PCR); OR
   - bDNA assay; OR
   - Tissue biopsy with pathological changes showing intra-nuclear inclusion bodies compatible with CMV infection (i.e. Owl’s eye)
   - Primary Infection - positive CMV IgM antibodies; OR
   - Reactivation - Positive CMV IgM antibodies with four-fold or greater increase in CMV IgG antibodies

2. Consolidation phase of treatment (maintenance phase post-induction with IV ganciclovir)

**Standard Approval Duration:** 3 to 6 months

Renewals will be considered for patients who continue to have active CMV infection. Renewal requests not meeting the criteria will be considered on a case-by-case basis but the physician must submit a rationale of why ongoing treatment is necessary.

**Standard Approval Duration:** 3 months

For **prophylaxis (prevention)** of **Epstein-Barr Virus (EBV)** infection in EBV D+/R- transplant recipients.

**Standard Approval Duration:** 6 months
**Valganciclovir**  
**Brand(s):** Valcyte  
**DOSAGE FORM/ STRENGTH:** 450 mg tablet, 50 mg/mL oral solution

For treatment of Epstein-Barr Virus (EBV) infection in transplant patients according to the following criteria:

1. Confirmed via biopsy (rare EBV positive cells present); OR

2. Objective evidence of active EBV infection (Patient must have one of the three below):
   - Newly positive or rising EBV PCR; OR
   - Reactivation: Positive anti-VCA IgM antibodies with a four-fold or greater increase in anti-VCA IgG antibodies; OR
   - Presence of EBV DNA or protein in pathologic tissue

**Standard Approval Duration:** 3 months

Notes:

Valganciclovir oral solution is considered for patients who meet the above requirements but who cannot swallow tablets or cannot use the tablets to achieve the planned dosing regimen.

Renewals for patients showing continued active infection with EBV PCR may be considered on a case-by-case basis by submitting the request to the EAP.

Requests for valganciclovir not meeting the above criteria or the Limited Use criteria on the formulary will be considered on a case-by-case basis but the prescriber must submit the request to the EAP.
Valganciclovir
Brand(s): Valcyte
DOSAGE FORM/STRENGTH: 450 mg tablet, 50 mg/mL oral solution

Approvals will be provided for the treatment of moderate to severe symptomatic congenital CMV (cCMV) in newborns who meet the following criteria:

- Prescribed by or in consultation with a pediatric ID specialist (from one of the 5 treatment centres in Ontario: London, Hamilton, Toronto, Kingston, Ottawa; or Winnipeg for the NorthWestern region of Ontario)
- Confirmed diagnosis of cCMV within the first 3 weeks¹ of birth by:
  - PCR (urine, saliva or quantitative serum CMV); OR
  - Positive culture results (urine or saliva)
- Treatment to start within one month of birth²
- Evidence of one or more of the following symptoms:
  - CNS disease (e.g., seizures, microcephaly, imaging abnormalities associated with CMV)
  - Eye disease (e.g., chorioretinitis)
  - Severe life-threatening organ dysfunction (must be described)³
- Regular monitoring of labs for toxicity while on therapy

Approval Duration: maximum 6 months at 16mg/kg/dose BID (with dose adjustments in renal dysfunction, < 32 weeks gestational age, etc.)

Renewals: No extensions will be provided unless extenuating circumstances for severely affected infants. Case-by-case review with rationale for continued treatment (must include pediatric ID specialist consult note)

Valganciclovir oral liquid will be approved for newborns.

All other requests not meeting the above criteria will be reviewed on a case-by-case basis including:
- Initiation of treatment after one month of age²
- Evidence of sensorineural hearing loss (SNHL) only (i.e., no other symptom described above)
- Isolated/multiple findings of mild symptoms such as: intrauterine growth retardation (IUGR), thrombocytopenia, elevated liver enzymes, jaundice, hepatitis

Sufficient rationale including consult from pediatric ID specialist must be provided before sending for external review.
Renewals of HIV Drugs

**Enfuvirtide**
*Brand(s):* Fuzeon  
*DOSAGE FORM/ STRENGTH:* 108 mg/vial Injection

Initial approvals require case-by-case review through the EAP upon receiving sufficient clinical information for an external review by a medical expert.

EAP will renew for patients who have responded to therapy and have undetectable viral load or increasing / stable CD4 count.

**Standard Approval Duration:** 6 months

**Tipranavir**
*Brand(s):* Aptivus  
*DOSAGE FORM/ STRENGTH:* 250 mg capsules

Initial approvals require case-by-case review through the EAP upon receiving sufficient clinical information for an external review by a medical expert.

EAP will renew for patients who have responded to therapy and have undetectable viral load or increasing / stable CD4 count.

**Standard Approval Duration:** 12 months
Renewals of Multiple Sclerosis Drugs

**Dimethyl Fumarate**  
Brand(s): Tecfidera  
**DOSAGE FORM/STRENGTH:** 120mg and 240 mg capsule

EAP will renew coverage of dimethyl fumarate for patients who are stable and experienced no more than one disabling attack/relapse in the past year and have an EDSS score less than or equal to 5.

Prescriber must provide the following information:

- Description of the patient's clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- EDSS score

**Dosage:** 120 mg twice daily  
**Maintenance:** 240 mg twice daily  

**Standard Approval Duration:** 2 years for first renewal, 5 years for 2nd and subsequent renewals

**Fingolimod**  
Brand(s): Gilenya and generics  
**DOSAGE FORM/STRENGTH:** 0.5 mg Capsule

EAP will renew coverage of Fingolimod for patients with RRMS who are stable and experienced no more than one disabling attack/relapse in the past year and have an EDSS score less than or equal to 5.5.

Prescriber must provide the following information:

- Description of the patient's clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- EDSS score
**Standard Approval Duration**: 2 years for first renewal, 5 years for 2nd and subsequent renewals

Renewal requests where patients have experienced more than 1 attack in the past year will be externally reviewed.

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**Glatiramer Acetate**  
*Brand(s):* Copaxone, Glatect  
*DOSAGE FORM/ STRENGTH:* 20 mg/mL Injection

All treatment naïve patients will be required to access Glatiramer biosimilar Glatect through the ODB formulary upon meeting Limited Use Criteria.

In RRMS/CDMS:

EAP will renew coverage of Glatiramer for patients who have benefited from therapy and have an EDSS score ≤ 6.

The physician must provide the following information:
- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- EDSS score

**Standard Approval Duration**: 2 years

In CIS:

EAP will renew coverage of Glatiramer only for patients who have benefited from therapy and have an EDSS score ≤ 6.

The physician must provide the following information:
- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- EDSS score

**Standard Approval Duration**: 2 years for first renewal, 5 years for 2nd and subsequent renewals

Renewal requests where patients have experienced more than 1 attack in the past year will be externally reviewed.
## Interferon beta-1a

**Brand(s):** Avonex, Rebif  
**DOSAGE FORM/STRENGTH:** (Avonex) 30 mcg/0.5 mL prefilled syringe, 30 mcgs prefilled autoinjector; (Rebif) 22 mcg and 44 mcg syringe injection, 66 mcg and 132 mcg prefilled cartridge

In RRMS/CDMS and CIS:

EAP will renew coverage of Interferon beta-1a only for patients who have benefited from therapy and have an EDSS score \( \leq 6 \).

The physician must provide the following information:
- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- EDSS score

**Standard Approval Duration:** 2 years for first renewal, 5 years for 2\(^{nd}\) and subsequent renewals

Renewal requests where patients have experienced more than 1 attack in the past year will be externally reviewed.

## Interferon beta-1b

**Brand(s):** Betaseron, Extavia  
**DOSAGE FORM/STRENGTH:** Betaseron 9.6 MIU = 0.3mg inj, Extavia 0.3 mg vial injection

IN RRMS/CDMS and CIS:

EAP will renew coverage of Interferon beta-1b only for patients who have benefited from therapy and have an EDSS score \( \leq 6 \).

The prescriber must provide the following information:
- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- EDSS score

**Standard Approval Duration:** 2 years for first renewal, 5 years for 2\(^{nd}\) and subsequent renewals

Renewal requests where patients have experienced more than 1 attack in the past year will be externally reviewed.
Natalizumab
Brand(s): Tysabri
DOSAGE FORM/ STRENGTH: 300 mg/15 mL

EAP will renew coverage of Natalizumab for patients with RRMS who have benefited from therapy and have an EDSS score less than or equal to 5.

The physician must provide the following information:
- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- EDSS score

Standard Approval Duration: 2 years

Standard Approval Duration: 2 years for first renewal, 5 years for 2nd and subsequent renewals

Renewal requests where patients have experienced more than 1 attack in the past year will be externally reviewed.

Ocrelizumab
Brand(s): Ocrevus
DOSAGE FORM/ STRENGTH: 14mg tablet

EAP will renew coverage of ocrelizumab for patients with Relapsed Refractory Multiple Sclerosis (RRMS) who are stable and experienced no more than one disabling attack/relapse in the past year and have an EDSS score less than or equal to 6.5.

Prescriber must provide the following information:
- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- EDSS score

Exclusion criteria:
1. Combination therapy with another disease modifying therapy for RRMS will not be reimbursed.
2. Patients with an EDSS score equal to or greater than 7.0

Standard Approval Duration: 18 months
Renewal requests where patients have experienced more than 1 attack in the past year will be externally reviewed.

**EAP will renew coverage of ocrelizumab for patients with Primary Progressive Multiple Sclerosis (PPMS) who continue to benefit from treatment and who have an Expanded Disability Status Scale (EDSS) score less than 7.0**

Prescriber must provide the following information:
- Description of the patient’s clinical status in the past year
- Date and details of the most recent neurological examination; and
- EDSS score

**Exclusion Criteria:**

Patients with an EDSS score equal to or greater than 7.0

Dosage: Initial dose of 300 mg intravenous infusion, followed 2 weeks later by a second 300 mg intravenous infusion. Subsequent doses of single 600 mg intravenous infusion every 6 months after the first initial dose.

Standard Approval Duration: 18 months

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**Peginterferon beta-1a**

**Brand(s): Plegridy**

**DOSAGE FORM/ STRENGTH:** 125mcg/0.5mL, 94mcg/0.5mL Injection, Starter Pack: 63mcg/0.5mL, 94mcg/0.5mL

EAP will renew coverage of peginterferon beta-1a only for patients who have benefited from therapy and have an EDSS score ≤ 6.

The prescriber must provide the following information:
- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- EDSS score

**Standard Approval Duration:** 2 years, 5 years for 2nd and subsequent renewals

Renewals

Renewal requests where patients have experienced more than 1 attack in the past year will be externally reviewed.
EAP will renew coverage of teriflunomide for patients who are stable and experienced no more than one disabling attack/relapse in the past year and have an EDSS score less than or equal to 5.

Prescriber must provide the following information:

- Description of the patient’s clinical course in the last year, including details of all attacks;
- Date and details of the most recent neurological examination (within the last 90 days); and
- EDSS score

Dosage: 14 mg daily

**Standard Approval Duration:** 2 years for first renewal, 5 years for 2\(^{nd}\) and subsequent renewals

Renewal requests where patients have experienced more than 1 attack in the past year will be externally reviewed.