

Request for Immunomodulatory Therapy (IMT) for Non-infectious Ocular Inflammatory Disease (OID): Exceptional Access Program (EAP)



To avoid delays, please ensure that all appropriate information for each section is provided.

| Section 1 – Physician Information | | | Section 2 – Patient Information | | |
|--|-------------|---|---|---------------------|-----------|
| First Name | Initial | Last Name | First Name | Initial | Last Name |
| Street # | Street Name | | OHIP Number | | |
| City | | Postal Code | Gender <input type="checkbox"/> Male <input type="checkbox"/> Female | Current Weight (kg) | |
| Fax | | Telephone (Back Line) | Date of Birth (DD/MM/YYYY) | | |
| Request Type <input type="checkbox"/> New Request (complete sections 3, 4 & 5) <input type="checkbox"/> Renewal Request (complete sections 3, 4 & 6) | | Is the patient currently taking the drug requested below? <input type="checkbox"/> Yes - Start Date (DD/MM/YYYY): <input type="checkbox"/> No | | | |
| | | EAP # | OR <input type="checkbox"/> TFA Mechanism Previously Used | | |

Section 3 – Drug, Dose and Regimen Requested

| | |
|---|------------------|
| <input type="checkbox"/> Mycophenolate Mofetil (Cellcept®) 1000 – 1500 mg po bid, or up to 1200 mg/m2/day divided BID for patient <18 years | Dosage |
| <input type="checkbox"/> Infliximab (Remicade®) 5-10 mg/kg IV at 0, 2, 6 weeks followed by maintenance therapy every 4-8 weeks | Dosing Frequency |
| <input type="checkbox"/> Adalimumab (Humira®) 40 mg subcutaneous every 1-2 weeks; For patients <18 years: 20 mg for patients <30 kg, 40 mg for patients ≥30 kg | |
| <input type="checkbox"/> Rituximab (Rituxan®)* Up to 1000 mg IV on days 1 and 15 and 3rd infusion at 6-12 months | |
| <input type="checkbox"/> Other (Specify dose, route and frequency of administration) : | |

*Rituximab is not funded for maintenance therapy. For subsequent rituximab requests following the initial requests, patients will be considered upon experiencing subsequent deterioration of symptoms at least 6 months from the last dose of rituximab.

Section 4 – Clinical Information

1. Specify the type ocular inflammatory disease (OID) for which the drug product is being requested:

| | |
|---|--|
| <input type="checkbox"/> Chronic Juvenile Idiopathic Arthritis (JIA) - associated uveitis | <input type="checkbox"/> Ocular mucous membrane pemphigoid |
| <input type="checkbox"/> Ocular inflammation associated with Behcet's disease | <input type="checkbox"/> Retinochoroidopathy |
| <input type="checkbox"/> Scleritis | <input type="checkbox"/> Serpiginous choroidopathy |
| <input type="checkbox"/> Uveitis | <input type="checkbox"/> Other (Specify): |

Specify if disease is: Anterior Intermediate Posterior Pan-Uveitis

2. Immediately vision-threatening OID? No Yes - Provide consultation notes/letter from specialist in OIDs to confirm severity

3. The OID affects: Left eye Right eye Both eyes

4. Specify the type of ocular specialist overseeing this patient's treatment:

Uveitis specialist Retinal specialist familiar with OIDs Pediatric ophthalmologist Other (Specify specialty):

Section 5 – Previous / Current Therapies

To avoid delays, provide details of ALL prior treatments with corticosteroids (include the route of administration) AND formulary immunosuppressants (e.g. methotrexate) or provide reasons why funded alternatives cannot be used.

| Name of Drug (Specify drug name) | Never prescribed | Route of administration (e.g. oral/topical/SC/intravitreal/subtenon) | Dose | Start Date (DD/MM/YYYY) | End Date (DD/MM/YYYY) | Response to therapy (e.g. efficacy, intolerance, etc.) / Contraindication and include reason for discontinuation |
|--|--------------------------|---|------|----------------------------|--------------------------|---|
| <input type="checkbox"/> Corticosteroid: | <input type="checkbox"/> | | | | | |
| <input type="checkbox"/> Methotrexate: | <input type="checkbox"/> | | | | | |
| <input type="checkbox"/> Other: | <input type="checkbox"/> | | | | | |
| <input type="checkbox"/> Other: | <input type="checkbox"/> | | | | | |
| <input type="checkbox"/> Other: | <input type="checkbox"/> | | | | | |

Section 6 – Renewal Information

Check all that apply and provide recent consultation notes/letter with clinical update and details of treatment response:

Improvement of vision Stability of vision Remission from/control of ocular inflammation

RITUXIMAB SPECIFICALLY, please check/complete all that apply and provide recent consultation notes/letter with clinical update:

Experiencing deterioration of symptoms

Infusion Date (DD/MM/YYYY): _____ Treatment response: _____

Infusion Date (DD/MM/YYYY): _____ Treatment response: _____

| | | |
|---------------------------------|-------------|-------------------|
| Physician Signature (Mandatory) | CPSO Number | Date (DD/MM/YYYY) |
|---------------------------------|-------------|-------------------|