

### **Prior Authorization Request**

**BENLYSTA** (belimumab)

#### Instructions

Please complete Part A and have your physician complete Part B. Completion and submission is not a guarantee of approval. Any fees related to the completion of this form are the responsibility of the plan member. Drugs in the Prior Authorization Program may be eligible for reimbursement if the patient does not qualify for coverage under a primary plan or a government program. Drugs used for indications not approved by Health Canada may be denied. For Quebec plan members, RAMQ exception drug criteria may apply. The decision for approval versus denial is based on pre-defined clinical criteria, primarily based on Health Canada approved indication(s) and on supporting evidence-based clinical protocols. The plan member will be notified whether their request has been approved or denied. Please note that you have the right to appeal the decision made by Express Scripts Canada.

Part A - Patient Patient information First Name: Last Name: Insurance Carrier Name/Number: Group Number: Client ID: Relationship: Employee Spouse Dependent Date of Birth (YYYY/MM/DD): Gender: Male Female Language: | English | French Address: City: Province: Postal Code: Email address: Telephone (home): Telephone (cell): Telephone (work): Coordination of benefits **Patient** Is the patient enrolled in any patient assistance program? Yes No **Assistance Program** Contact Name: \_ Has the patient applied for reimbursement under a provincial plan? Yes No N/A **Provincial** Coverage What is the coverage decision of the drug? Approved Denied \*Attach decision letter\* Has the patient applied for reimbursement under a primary plan? Yes No N/A **Primary** Coverage What is the coverage decision of the drug? Approved Denied \*Attach decision letter\* **Authorization** 

On behalf of myself and my eligible dependents, I authorize my group benefit provider, and its agents, to exchange the personal information contained on this form. I give my consent on the understanding that the information will be used solely for purposes of administration and management of my group benefit plan. This consent shall continue so long as my dependents and I are covered by, or are claiming benefits under the present group contract, or any modification, renewal, or reinstatement thereof.

| Plan Member Signature | <br>Date |
|-----------------------|----------|



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#### Part B - Prescriber

Please see instructions on page 1 and complete all sections below. <u>Incomplete forms may result in automatic denial</u>. Please do **not** provide genetic test information or results.

## SECTION 1 - DRUG REQUESTED BENLYSTA (belimumab) New request Renewal request\* Dose Administration (ex: oral, IV, etc) Duration Frequency Site of drug administration: Physician's office/Infusion clinic Hospital (inpatient) Home Hospital (outpatient) \* Please submit proof of prior coverage if available SECTION 2 - ELIGIBILITY CRITERIA 1. Please indicate if the patient satisfies the below criteria: Systemic Lupus Erythematosus For the treatment of systemic lupus erythematosus (SLE) in an adult, AND The patient has autoantibody (ANA or dsDNA) positive results, AND The patient has a Safety of Estrogens in Lupus National Assessment-Systemic Lupus Erythematosus Disease Activity Index (SELENA-SLEDAI) score of 6 or greater, AND The patient has had an inadequate response or has a documented intolerance to corticosteroids (Please list prior therapies in the chart below), AND The patient has had an inadequate response or has a documented intolerance to either an antimalarial (e.g. hydroxychloroquine), or an immunosuppressant (e.g. methotrexate) (*Please list prior therapies in the chart below*) **Active Lupus Nephritis** For the treatment of active lupus nephritis in an adult, AND The patient has received standard therapy with corticosteroids and mycophenolate mofetil for induction and maintenance therapy, OR The patient has received standard therapy with corticosteroids and cyclophosphamide for induction therapy and azathioprine for maintenance therapy, AND The patient has not received dialysis in the past 12 months OR None of the above criteria applies. Relevant additional information:



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| Drug | Decede and                | Duration of therapy |    | Reason for cessation |                         |
|------|---------------------------|---------------------|----|----------------------|-------------------------|
|      | Dosage and administration | From                | То | Inadequate response  | Allergy/<br>Intolerance |
|      |                           |                     |    |                      |                         |
|      |                           |                     |    |                      |                         |
|      |                           |                     |    |                      |                         |
|      |                           |                     |    |                      |                         |
|      |                           |                     |    |                      |                         |
|      |                           |                     |    |                      |                         |

#### **SECTION 3 - PRESCRIBER INFORMATION**

| Physician's Name:    |            |
|----------------------|------------|
| Address:             |            |
| Tel:                 | Fax:       |
| License No.:         | Specialty: |
| Physician Signature: | Date:      |

Please fax or mail the completed form to Express Scripts Canada®

**Fax:** Express Scripts Canada Clinical Services 1 (855) 712-6329

**Mail:** Express Scripts Canada Clinical Services 5770 Hurontario Street, 10<sup>th</sup> Floor Mississauga, ON L5R 3G5