



Benlysta (belimumab)

Fax completed form to: (855) 840-1678
If this is an URGENT request, please call (800) 882-4462

PHYSICIAN INFORMATION			PATIENT INFORMATION		
* Physician Name:			*Due to privacy regulations we will not be able to respond via fax with the outcome of our review unless all asterisked (*) items on this form are completed.*		
Specialty:	* DEA, NPI or TIN:				
Office Contact Person:			* Patient Name:		
Office Phone:			* Cigna ID:	* Date of Birth:	
Office Fax:			* Patient Street Address:		
Office Street Address:			City:	State:	Zip:
City:	State:	Zip:	Patient Phone:		

Urgency: Standard Urgent (In checking this box, I attest to the fact that applying the standard review time frame may seriously jeopardize the customer's life, health, or ability to regain maximum function)**Medication requested:** Benlysta 120mg vial Benlysta 400mg vial

ICD10:

Directions for use:

Quantity:

Duration of therapy:

J-Code:

Dose (in Mg/kg):

Frequency (for example Weeks 0, 2):

Will the dose be administered at Weeks 0, 2, and 4, with subsequent doses separated by at least 4 weeks?

Yes No **Where will this medication be obtained?** Accredo Specialty Pharmacy** Hospital Outpatient Retail pharmacy Other (please specify): Home Health / Home Infusion vendor Physician's office stock (billing on a medical claim form)****Cigna's nationally preferred specialty pharmacy**

****Medication orders can be placed with Accredo via E-prescribe - Accredo (1620 Century Center Pkwy, Memphis, TN 38134-8822 | NCPDP 4436920), Fax 888.302.1028, or Verbal 866.759.1557**

Facility and/or doctor dispensing and administering medication:

Facility Name:

State:

Tax ID#:

Address (City, State, Zip Code):

Where will this drug be administered? Patient's Home Hospital Outpatient Physician's Office Other (please specify):

NOTE: Per some Cigna plans, infusion of medication MUST occur in the least intensive, medically appropriate setting.

Is this patient a candidate for re-direction to an alternate setting (such as alternate infusion site, physician's office, home) with assistance of a Specialty Care Options Case Manager? Yes No (provide medical necessity rationale):

Is the requested medication for a chronic or long-term condition for which the prescription medication may be necessary for the life of the patient? Yes No

Diagnosis:

- Lupus Nephritis (LN)
 Rheumatoid Arthritis
 Systemic Lupus Erythematosus (SLE)
 Other: (Please provide the patient's diagnosis or reason for treatment):

Clinical Information:

Will the requested medication be used in combination with another biologic agent (for example, Cimzia, an etanercept product [Enbrel, biosimilars], an adalimumab product [Humira, biosimilars], Saphnelo, Simponi [Aria or SC], an infliximab product [Remicade, biosimilars], Actemra [SC or IV], Kevzara, Orencia [SC or IV], a rituximab product [Rituxan, biosimilars], Kineret, Stelara [SC or IV], Siliq, Cosentyx, Taltz, Ilumya, Tremfya, Skyrizi, Entyvio, Zymfentra, Omvoh, and Bimzelx)? Yes No

Is the requested medication being used in combination with Lupkynis (voclosporin capsules)? Yes No

If LN
 Is this medication being prescribed by, or in consultation with, a nephrologist or rheumatologist? Yes No

Is the requested medication being used concurrently with an immunosuppressive regimen? Note: Examples of an immunosuppressive regimen include azathioprine, cyclophosphamide, leflunomide, methotrexate, mycophenolate mofetil, and/or a systemic corticosteroid. Yes No

Is the patient currently receiving Benlysta intravenous or subcutaneous? Yes No

(if no) Has the diagnosis of lupus nephritis been confirmed on biopsy? Please Note: For example, World Health Organization class III, IV, or V lupus nephritis. Yes No

(if yes) According to the prescriber, has the patient responded to Benlysta subcutaneous or intravenous? Please Note: Examples of a response include improvement in organ dysfunction, reduction in flares, reduction in corticosteroid dose, decrease of anti-dsDNA titer, and improvement in complement levels (that is, C3, C4). Yes No

If SLE
 Is this medication being prescribed by or in consultation with a rheumatologist, clinical immunologist, nephrologist, neurologist, or dermatologist? Yes No

Is the requested medication being used concurrently with at least ONE other standard therapy for systemic lupus erythematosus (SLE)? Note: Examples of standard therapies include an antimalarial (for example, hydroxychloroquine), systemic corticosteroid (for example, prednisone), and other immunosuppressants (for example, azathioprine, mycophenolate mofetil, methotrexate). Yes No

(if no) According to the prescriber, is the patient intolerant to standard therapy due to a significant toxicity? Yes No

Is the patient currently receiving Benlysta intravenous or subcutaneous? Yes No

(if no) Does the patient have autoantibody-positive systemic lupus erythematosus (SLE), defined as positive for antinuclear antibodies (ANA) and/or anti-double-stranded DNA (anti-dsDNA) antibody? Note: Not all patients with SLE are positive for anti-dsDNA, but most will be positive for ANA. Yes No

(if yes) According to the prescriber, has the patient responded to Benlysta subcutaneous or intravenous? Please Note: Examples of a response include reduction in flares, reduction in corticosteroid dose, decrease of anti-dsDNA titer, improvement in complement levels (that is, C3, C4), or improvement in specific organ dysfunction (for example, musculoskeletal, blood, hematologic, vascular, others). Yes No

Additional pertinent information: *Please provide any additional pertinent clinical information*

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Prescriber Signature: _____ **Date:** _____

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