

**Pharmacy Prior Approval Request for  
Lupus Medications**

**Beneficiary Information**

1. Beneficiary Last Name: \_\_\_\_\_ 2. First Name: \_\_\_\_\_  
3. Beneficiary ID #: \_\_\_\_\_ 4. Beneficiary Date of Birth: \_\_\_\_\_ 5. Beneficiary Gender: \_\_\_\_\_

**Prescriber Information**

6. Prescribing Provider NPI #: \_\_\_\_\_  
7. Requester Contact Information - Name: \_\_\_\_\_ Phone #: \_\_\_\_\_ Ext. \_\_\_\_\_

**Drug Information**

8. Drug Name: \_\_\_\_\_ 9. Strength: \_\_\_\_\_ 10. Quantity Per 30 Days: \_\_\_\_\_  
11. Length of Therapy (in days):  up to 30 Days  60 Days  90 Days  120 Days  180 Days  365 Days

**Clinical Information****Initial authorization (answer questions 1-10)**

1. Is the medication being prescribed by or in consultation with a rheumatologist or nephrologist?  **Yes**  **No**
2. Is the member  $\geq 18$  years of age or older?  **Yes**  **No**
3. Does the member have a diagnosis of active systemic lupus erythematosus (SLE)?  **Yes**  **No**
4. Is the member auto-antibody positive?  **Yes**  **No**
5. Is the member utilizing the medicine in combination with standard treatment regimens (NSAIDs, corticosteroids, anti-malarials, or immunosuppressive drugs) or standard treatment regimens were not tolerated or beneficial?  **Yes**  **No**
6. Is the medication being used concurrently with other biologics and/or IV cyclophosphamide?  **Yes**  **No**
7. Does the member have a diagnosis or severe active central nervous system lupus?  **Yes**  **No**

**OR**

8. Does the member have a diagnosis of active lupus nephritis?  **Yes**  **No**
9. Is the member utilizing the medicine in combination with standard treatment regimens (immunosuppressive therapy) with the exception of cyclophosphamide?  **Yes**  **No**
10. Does the member have baseline blood pressure  $\leq 165/105$  mg Hg, estimated glomerular filtration rate is  $\geq 45$  mL/min/1.73 m<sup>2</sup>

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**Pharmacy Prior Approval Request for****For re-authorization (answer questions below)**

**11.** Is there documented improvement in functional impairment such as fewer flares that required steroid treatment, lower average daily oral prednisone dose, improved daily function either as measured through a validated functional scale or through improved daily performance documented at clinic visits, or sustained improvement in laboratory measures of lupus activity?  Yes  No

**12.** Is there documentation of any treatment restricting adverse effects (e.g. hypertension, neurotoxicities, hyperkalemia)

**\*\*Please attach current progress notes documenting disease status and clinical response to the medicine.\*\***

Signature of Prescriber: \_\_\_\_\_ Date: \_\_\_\_\_

**(Prescriber Signature Mandatory)**

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.

Fax this form to: (866) 399-0929

Pharmacy PA Call Center: (833) 585-4309