

Field Name	Field Description
Prior Authorization Group Description	<b>Benlysta (belimumab)</b>
Drugs	Benlysta (belimumab)
Covered Uses	Medically accepted indications are defined using the following sources: the Food and Drug Administration (FDA), Micromedex, the Drug Package Insert, and/or per the standard of care guidelines
Exclusion Criteria	Severe active central nervous system lupus
Required Medical Information	See “other criteria”
Age Restrictions	Must be at least 5 years of age
Prescriber Restrictions	Prescribed by or in consultation with a rheumatologist or nephrologist
Coverage Duration	If all the criteria are met initial authorization requests may be approved for up to 6 months. Reauthorization requests may be approved for up to 12 months.
Other Criteria	<p><b><u>Initial Authorization:</u></b></p> <ul style="list-style-type: none"> <li>• <u>Active systemic lupus erythematosus (SLE)</u> <ul style="list-style-type: none"> <li>○ Provider attestation that the patient is positive for autoantibodies (or antinuclear antibodies or anti–double-stranded DNA [anti-dsDNA] antibodies)</li> <li>○ The member has tried and failed both of the following (or contraindication/inability to use these medications): <ul style="list-style-type: none"> <li>▪ Hydroxychloroquine</li> <li>▪ One other immunosuppressant [e.g., methotrexate, azathioprine, calcineurin inhibitors or mycophenolate]</li> </ul> </li> </ul> </li> <li>• <u>Active lupus nephritis</u> <ul style="list-style-type: none"> <li>○ Provider attestation of diagnosis confirmed by kidney biopsy</li> <li>○ The member has tried and failed, or has a medical reason for not using, both of the following <ul style="list-style-type: none"> <li>▪ Cyclophosphamide or tacrolimus</li> <li>▪ Mycophenolate</li> </ul> </li> </ul> </li> <li>• Provider states the member will not be receiving concomitant therapy with the following: <ul style="list-style-type: none"> <li>○ B-cell targeted therapy including (but not limited to) rituximab</li> <li>○ Interferon receptor antagonist, type 1 including (but not limited to) Saphnelo (anifrolumab)</li> </ul> </li> <li>• Dosing is appropriate per labeling</li> </ul> <p><b><u>Criteria for Reauthorization:</u></b></p> <ul style="list-style-type: none"> <li>• Documentation or provider attestation of positive clinical response as indicated by one of the following:</li> </ul>

<p>Revision/Review Date: 1/2023</p>	<ul style="list-style-type: none"> <li>○ Fewer flares that required steroid treatment</li> <li>○ Lower average daily oral prednisone dose</li> <li>○ Improved daily function either as measured through a validated functional scale or through improved daily performance documented at clinic visits</li> <li>○ Sustained improvement in laboratory measures of lupus activity</li> <li>● Dosing is appropriate per labeling</li> </ul> <p><b>Medical Director/clinical reviewer must override criteria when, in his/her professional judgement, the requested item is medically necessary.</b></p>
---	---