

Field Name	Field Description
Prior Authorization Group Description	Benlysta (belimumab)
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><u>Initial Authorization:</u></p> <ul style="list-style-type: none"> • <u>Active systemic lupus erythematosus (SLE)</u> <ul style="list-style-type: none"> ○ Provider attestation that the patient is positive for autoantibodies (or antinuclear antibodies or anti–double-stranded DNA [anti-dsDNA] antibodies) ○ The member has tried and failed both of the following (or contraindication/inability to use these medications): <ul style="list-style-type: none"> ▪ Hydroxychloroquine ▪ One other immunosuppressant [e.g., methotrexate, azathioprine, calcineurin inhibitors or mycophenolate] • <u>Active lupus nephritis</u> <ul style="list-style-type: none"> ○ Provider attestation of diagnosis confirmed by kidney biopsy ○ The member has tried and failed, or has a medical reason for not using, both of the following <ul style="list-style-type: none"> ▪ Cyclophosphamide or tacrolimus ▪ Mycophenolate • Provider states the member will not be receiving concomitant therapy with the following: <ul style="list-style-type: none"> ○ B-cell targeted therapy including (but not limited to) rituximab ○ Interferon receptor antagonist, type 1 including (but not limited to) Saphnelo (anifrolumab) • Dosing is appropriate per labeling <p><u>Criteria for Reauthorization:</u></p> <ul style="list-style-type: none"> • Documentation or provider attestation of positive clinical response as indicated by one of the following: <ul style="list-style-type: none"> ○ Fewer flares that required steroid treatment

<p>Revision/Review Date: 2/2025</p>	<ul style="list-style-type: none"> ○ 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 ○ Sustained improvement in laboratory measures of lupus activity <ul style="list-style-type: none"> ● Dosing is appropriate per labeling <p>Medical Director/clinical reviewer must override criteria when, in his/her professional judgement, the requested item is medically necessary.</p>
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