



Benlysta[®] (belimumab) (Intravenous)

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I. Length of Authorization

Coverage will be provided for 12 months and may be renewed.

II. Dosing Limits

A. Quantity Limit (max daily dose) [NDC Unit]:

- Loading Dose (doses administered on days 1, 15 and 29):
 - \circ Benlysta 120 mg single-dose vial for injection: 9 vials per 29 days
 - \circ Benlysta 400 mg single-dose vial for injection: 9 vials per 29 days
- Maintenance Dose:
 - \circ $\;$ Benlysta 120 mg single-dose vial for injection: 3 vials per 28 days
 - o Benlysta 400 mg single-dose vial for injection: 3 vials per 28 days

B. Max Units (per dose and over time) [HCPCS Unit]:

- Loading Dose:
 - \circ 120 billable units on days 1, 15 and 29
- Maintenance Dose:
 - \circ 120 billable units per 28 days

III. Initial Approval Criteria¹

- For patients age 18 years and older, patient must have tried and failed, or have a documented contraindication to, subcutaneous Benlysta before the intravenous infusion will be authorized.
- Patient is at least 5 years of age; AND

Universal Criteria¹

• Patient must not have an active infection; AND



- Patient will not receive live vaccines during therapy or within 30 days prior to starting treatment; **AND**
- Patient does not have severe active central nervous system lupus; AND
- Will be used in combination with standard therapy (e.g., anti-malarials, corticosteroids, non-steroidal anti-inflammatory drugs, immunosuppressives); **AND**
- Will not be used in combination with rituximab, anifrolumab, or IV cyclophosphamide*; **AND**
- * Note: May be used in combination with IV cyclophosphamide for Lupus Nephritis.

Systemic Lupus Erythematosus (SLE) † 1,9,11,12, 22-27,29

- Patient has a confirmed diagnosis of SLE as evidenced by <u>all</u> of the following:
 - Confirmed SLE classification criteria score ≥10* (*Note: must include clinical and immunologic domains criteria*)
 - Anti-nuclear antibody (ANA) titer of ≥1:80 measured via indirect immunofluorescence (IIF) on human epithelial (HEp-2) cells (or an equivalent ANA positive test) at least once; AND
- Patient has documented active disease; AND
- Physician has assessed baseline disease severity utilizing an objective measure/tool (i.e., Systemic Lupus Erythematosus Disease Activity Index 2000 (SLEDAI 2K), British Isles Lupus Assessment Group-2004 (BILAG 2004), and/or Physician's Global Assessment (PGA) score); **AND**
- Patient has failed to respond adequately to at least one (1) standard therapy such as antimalarials, corticosteroids, non-steroidal anti-inflammatory drugs, immunosuppressives**

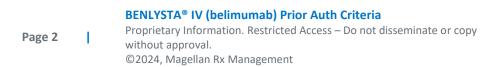
**** Note:** For patients already established on biologic therapy, trial and failure of standard therapy is not required.

Lupus Nephritis (LN) † ^{1,9,11,12,19,22,28-30}

- Patient has active lupus nephritis Class III, IV, or V as confirmed by renal biopsy; AND
- Patient has a confirmed diagnosis of SLE as evidenced by <u>all</u> of the following**:
 - Confirmed SLE classification criteria score ≥10* (*Note: must include clinical and immunologic domains criteria*)
 - Anti-nuclear antibody (ANA) titer of ≥1:80 measured via indirect immunofluorescence (IIF) on human epithelial (HEp-2) cells (or an equivalent ANA positive test) at least once; AND
- Physician has obtained a baseline measurement of one or more of the following: urine protein:creatinine ratio (uPCR), estimated glomerular filtration rate (eGFR), or urine protein

******Patients with class III, IV, or V disease that do not meet the SLE diagnostic criteria will be reviewed on a case-by-case basis

*Classification Criteria for Systemic Lupus Erythematosus (SLE) ²²





Clinical Score ^A (range: 0-39)	Clinical Domains and Criteria	
-	Ormetitetionel: Use also al france 101°E	
2	Constitutional: Unexplained fever > 101°F	
	Hematologic:	
3	White blood cell count < 4,000/mm ³	
4	Platelet count < 100,000/mm ³ or Autoimmune hemolysis	
	Neuropsychiatric:	
2	Delirium	
3	Psychosis	
5	Primary generalized seizure or partial/focal seizure	
	Mucocutaneous +:	
2	Non-scarring alopecia or oral ulcers	
4	Subacute cutaneous or discoid lupus	
6	Acute cutaneous lupus	
	Serosal:	
5	Pleural or pericardial effusion	
6	Acute pericarditis	
	Musculoskeletal:	
6	Joint involvement with either synovitis involving 2 or more joints	
	with swelling or effusion OR tenderness in 2 or more joints with at	
	least 30 minutes of morning stiffness	
	Renal:	
4	Proteinuria > 0.5g/24 hr by a 24-hour urine or equivalent spot urine	
	protein-to-creatinine ratio	
8	Renal biopsy class II or V lupus nephritis	
10	Renal biopsy Class III or IV lupus nephritis	
Immunologic Score △ (range: 0-12)	Immunologic Domains and Criteria	
2	Presence of antiphospholipid antibodies (i.e., positive lupus	
	anticoagulant, positive anti-B2GP1 antibodies, and/or anti-	
	cardiolipin antibodies at medium or high titer)	
	Presence of low complement proteins (below lower limit of normal):	
3	Low C3 OR low C4	
4	Low C3 AND C4	
6	Presence of anti-Sm and/or anti-dsDNA antibodies	
*A web-based scoring calculator as well as further definitions of each criterion are available at:		

https://rheumatology.org/criteria

^AOccurrence on at least one occasion is sufficient to count toward score when all other causes have been ruled out. Count only the highest weighted score within each of the 10 domains (7 clinical and 3 immunologic) and any additional criteria within the same domain will not count. + Observed by a physician via clinical exam or photograph review

 \dagger FDA Approved Indication(s); \ddagger Compendia Recommended Indication(s); Φ Orphan Drug

IV. Renewal Criteria¹

Coverage can be renewed based on the following criteria:

• Patient continues to meet universal and other indication-specific relevant criteria such as concomitant therapy requirements (not including prerequisite therapy), performance status, etc. identified in section III; **AND**



• Absence of unacceptable toxicity from the drug. Examples of unacceptable toxicity include: depression, suicidal thoughts, serious infections, suspected or confirmed progressive multifocal leukoencephalopathy (PML), malignancy, severe hypersensitivity reactions/anaphylaxis, serious infusion-related reactions, etc.; **AND**

Systemic Lupus Erythematosus (SLE) 1,9,21-27,29

- Adequate documentation of disease stability and/or improvement as indicated by one or more of the following when compared to pre-treatment baseline:
 - Improvement in the SELENA-SLEDAI-2K; OR
 - Reduction of baseline BILAG-2004 (e.g. reduction from A to B or from B to C/D, and no BILAG-2004 worsening in other organ systems, as defined by ≥2 new BILAG-2004 B or ≥1 new BILAG A); OR
 - No worsening (<0.30-point increase) in Physician's Global Assessment (PGA) score; OR
 - Seroconverted (negative)

Lupus Nephritis (LN) 1,19,22,30

- Adequate documentation of disease stability and/or improvement as indicated by one or more of the following when compared to pre-treatment baseline:
 - Urine protein:creatinine ratio (uPCR); OR
 - Estimated glomerular filtration rate (eGFR); OR
 - Urine protein

V. Dosage/Administration¹

Indication	Dose
Systemic Lupus Erythematosus	• Loading Dose: 10 mg/kg intravenously (by a healthcare provider) every 2 weeks for the first 3 doses (days 1, 15 and 29)
(SLE) or Lupus Nephritis (LN)	• Maintenance Dose: 10 mg/kg intravenously (by a healthcare provider) every 4 weeks

VI. Billing Code/Availability Information

HCPCS Code:

• J0490 – Injection, belimumab, 10 mg; 1 billable unit = 10 mg

NDC(s):

- Benlysta 120 mg/5 mL single-dose vial for injection: 49401-0101-xx
- Benlysta 400 mg/20 mL single-dose vial for injection: 49401-0102-xx

VII. References

1. Benlysta [package insert]. Philadelphia, PA; GlaxoSmithKline LLC; February 2024. Accessed February 2024.

		BENLYSTA [®] IV (belimumab) Prior Auth Criteria
Page 4	1.1	Proprietary Information. Restricted Access – Do not disseminate or copy
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ICD-10	ICD-10 Description	
M32.10	Systemic lupus erythematosus organ or system involvement unspecified	
M32.11	Endocarditis in systemic lupus erythematosus	
M32.12	Pericarditis in systemic lupus erythematosus	
M32.13	Lung involvement in systemic lupus erythematosus	
M32.14	Glomerular disease in systemic lupus erythematosus	
M32.15	Tubulo-interstitial nephropathy in systemic lupus erythematosus	
M32.19	Other organ or system involvement in systemic lupus erythematosus	
M32.8	Other forms of systemic lupus erythematosus	
M32.9	Systemic lupus erythematosus, unspecified	

Appendix 1 – Covered Diagnosis Codes

Appendix 2 – Centers for Medicare and Medicaid Services (CMS)

The preceding information is intended for non-Medicare coverage determinations. Medicare coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determinations (NCDs) and/or Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. Local Coverage Articles (LCAs) may also exist for claims payment purposes or to clarify benefit eligibility under Part B for drugs which may be self-administered. The following link may be used to search for NCD, LCD, or LCA documents: https://www.cms.gov/medicare-coverage-database/search.aspx. Additional indications, including any preceding information, may be applied at the discretion of the health plan.

Medicare Part B Administrative Contractor (MAC) Jurisdictions				
Jurisdiction	Applicable State/US Territory	Contractor		
E (1)	CA, HI, NV, AS, GU, CNMI	Noridian Healthcare Solutions, LLC		
F (2 & 3)	AK, WA, OR, ID, ND, SD, MT, WY, UT, AZ	Noridian Healthcare Solutions, LLC		
5	KS, NE, IA, MO	Wisconsin Physicians Service Insurance Corp (WPS)		

Medicare Part B Covered Diagnosis Codes (applicable to existing NCD/LCD/LCA): N/A



	Medicare Part B Administrative Contractor (MAC) Jurisdictions				
Jurisdiction	Applicable State/US Territory	Contractor			
6	MN, WI, IL	National Government Services, Inc. (NGS)			
H (4 & 7)	LA, AR, MS, TX, OK, CO, NM	Novitas Solutions, Inc.			
8	MI, IN	Wisconsin Physicians Service Insurance Corp (WPS)			
N (9)	FL, PR, VI	First Coast Service Options, Inc.			
J (10)	TN, GA, AL	Palmetto GBA			
M (11)	NC, SC, WV, VA (excluding below)	Palmetto GBA			
L (12)	DE, MD, PA, NJ, DC (includes Arlington & Fairfax counties and the city of Alexandria in VA)	Novitas Solutions, Inc.			
K (13 & 14)	NY, CT, MA, RI, VT, ME, NH	National Government Services, Inc. (NGS)			
15	КҮ, ОН	CGS Administrators, LLC			

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