AvMed

MEDICAL PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions:</u> The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; <u>fax to 1-877-535-1391</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If information provided is not complete, correct, or legible, authorization can be delayed</u>.

For Medicare Members: 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 Determination (NCD) and Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. They can be found at: https://www.cms.gov/medicare-coverage-database/overview-and-quick-search.aspx. Additional indications may be covered at the discretion of the health plan.

Drug Requested: Saphnelo[™] (anifrolumab) for IV Infusion (J0491) (Medical)

MEMBER & PRESCRIBER IN	FORMATION: Authorization may be delayed if incomplete.	
Member Name:		
Member AvMed #:		
Prescriber Name:		
	Date:	
Office Contact Name:		
	Fax Number:	
DEA OR NPI #:		
Drug Form/Strength:		
Dosing Schedule:	Length of Therapy:	
Diagnosis:	ICD Code, if applicable:	
Weight:	Date:	
	ox, the timeframe does not jeopardize the life or health of the member of the function and would not subject the member to severe pain.	
Recommended Dosage: 300 mg ev	very 4 weeks	

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support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied. (Trials will be verified using pharmacy claims and/or submitted

chart notes.)

_ D	iagnosis: Moderate-to-Severe	Systemic Lupus Erythemato	sus (SLE)	
Initi	al Authorization: 12 months			
	Prescribed by or in consultation with	a rheumatologist or nephrologist		
	Member is 18 years of age or older			
	Member has a diagnosis of autoantib results for documentation):	ody-positive SLE confirmed by ON	NE of the following (submit lab	
	☐ anti-nuclear antibody (ANA) tit	$er \ge 1:80$		
	☐ anti-double stranded DNA (anti-	$-dsDNA) \ge 30 \text{ IU/mL}$		
	□ anti-Smith (anti-SM) antibody l	evels elevated according to reference	ce range	
	Member has active moderate to seve results for documentation):	re SLE activity as confirmed by ON	NE of the following (submit	
	☐ Systemic Lupus Erythematosus I	Disease Activity Index 2000 (SLED	AI-2K) score of ≥ 6	
	☐ British Isles Lupus Assessment C	Group (BILAG) 2004 organ domain	score of $\geq 1A$ or $\geq 2B$	
☐ Member has tried <u>THREE</u> of the following (verified by chart notes or pharmacy paid claims; check all that apply):				
	□ mycophenolate	□ hydroxychloroquine	□ azathioprine	
	□ cyclophosphamide	□ methotrexate	□ cyclosporine	
	□ corticosteroids	□ Other:		
	☐ Member is currently established on at least <u>ONE</u> of the following therapies taken for the last 90 days and will continue on current therapy if approved for Saphnelo (verified by chart notes or pharmacy paid claims; check all that apply):			
	□ mycophenolate	hydroxychloroquine	□ azathioprine	
	□ cyclophosphamide	□ methotrexate	□ cyclosporine	
	□ corticosteroids	□ Other:		
	Saphnelo [™] will NOT be approved to	for members with any of the follow	ing:	
	• Severe active central nervous sy	stem lupus		
	• Severe active lupus nephritis			
	 Concurrent use with cyclophosp belimumab (Benlysta) 	hamide, voclosporin or other biolog	gic therapies, including	
□ N	Ioderate-to-Severe Systemic L	upus Erythematosus (SLE)		

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Reauthorization Approval: 12 months. Check below all that apply. All criteria must be met for approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart

notes, must be provided or request may be denied.

	Member has experienced a positive clinical response to Saphnelo TM therapy as confirmed by <u>ONE</u> of the following (submit results for documentation):		
	 Reduction of all baseline BILAG A to B/C/D and baseline BILAG B to C/D, and no BILAG worsening in other organ systems, as defined by ≥ 1 new BILAG A or ≥ 2 new BILAG B 		
	□ No worsening from baseline in SLEDAI-2K, where worsening is defined as an increase from baseline of > 0 points in SLEDAI-2K		
	Member has an absence of intolerable side effects such as serious or recurrent infections, malignancy, severe hypersensitivity reactions/anaphylaxis or intolerable infusion reactions		
Med	dication being provided by (check box below that applies):		
	Location/site of drug administration:		
NPI or DEA # of administering location:			
	OR		
	Specialty Pharmacy - PropriumRx		
review	gent reviews: Practitioner should call AvMed Pre-Authorization Department if they believe a standard would subject the member to adverse health consequences. AvMed's definition of urgent is a lack of ent that could seriously jeopardize the life or health of the member or the member's ability to regain		

**Use of samples to initiate therapy does not meet step edit/ preauthorization criteria. **

*Previous therapies will be verified through pharmacy paid claims or submitted chart notes. *

maximum function.