HEALTHY U CHIP

PRIOR AUTHORIZATION REQUEST FORM SAPHNELO™

For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U CHIP Prior Authorization Department at 801-213-1547. Failure to submit clinical documentation to support this request will result in a dismissal of the request. If you have medical pharmacy prior authorization questions, please call for assistance: 833-981-0212 Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements. ID#: Date: Member Name: Gender: Physician: DOB: Office Phone: Office Fax: Office Contact: Height/Weight: HCPCS Code: Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria. **Product being requested:** □ Saphnelo[™] (anifrolumab-fnia) Dosing/Frequency: If the request is for reauthorization, proceed to reauthorization section. Questions Yes No **Comments/Notes MODERATE TO SEVERE SYSTEMIC LUPUS ERYTHEMATOSUS (SLE)** Please provide documentation 1. Does the member have a diagnosis of moderate to severe systemic lupus erythematosus (SLE)? 2. Have laboratory tests been completed indicating the presence Please provide documentation of autoantibodies (ANA, Anti-dsDNA, Anti-Sm, Anti-Ro/SSA, Anti-La/SSB)? 3. Is the requesting provider a rheumatologist or in consultation Please provide documentation with a rheumatologist? 4. Does the member have a Systemic Lupus Erythematosus Please provide documentation Disease Activity Index 2000 (SLEDAI-2K) score of \geq 6 points? 5. Does the member have a contraindication, intolerance or Please provide documentation failure to Benlysta®? 6. Is the member receiving Saphnelo™ in combination with a Please provide documentation biologic agent, Benlysta® or cyclophosamide?

Please provide documentation

7. Does the member have active musculoskeletal or cutaneous

glucocorticoids, antimalarials and/or other immunosuppressive

disease that is unresponsive to standard therapy with

agents?

 8. Does the member have corticosteroid-dependent disease (prednisone equivalent dose ≥7.5mg/day) or trialed and failed both of the following: hydroxychloroquine AND at least 1 immunosuppressant (e.g., azathioprine, methotrexate, mycophenolate, cyclosporine, cyclophosphamide) 			Please provide documentation
9. Is the member at least 80% compliant for at least 6 months with their baseline therapy (i.e. glucocorticoids, immunosuppressants and/or antimalarials)?			Please provide documentation
10. Will the member use Saphnelo™ concurrently with baseline therapy, unless the member has a contraindication or intolerance to all?			Please provide documentation
11.Does the member have severe active lupus nephritis or severe active central nervous system lupus (e.g., generalized seizures, psychosis, stroke, peripheral neuropathies)?			Please provide documentation
REAUTHORIZATION			
1. Is the request for reauthorization of therapy?			Place provide decumentation
2. Has the member shown a positive clinical response to therapy?			Please provide documentation
3. Has the member been compliant with baseline therapy during Saphnelo™ administration, unless otherwise contraindicated?			Please provide documentation
4. Is the member receiving Saphnelo™ in combination with a biologic agent or Benlysta®?			Please provide documentation
5. Does the member have severe active central nervous system lupus or severe active lupus nephritis?			Please provide documentation
What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc. Additional information:			
Physician Signature:			

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Policy: PHARM-CHIP-M037 Origination Date: 07/01/2024 Reviewed/Revised Date: Next Review Date:

Current Effective Date: 07/01/2024

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