## SENTARA HEALTH PLANS

## PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST\*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request.</u> All other information may be filled in by office staff; <u>fax to 1-800-750-9692</u>. No additional phone calls will be necessary if all information (including phone and fax #s) on this form is correct. <u>If the information provided is not complete, correct, or legible, the authorization process may be delayed.</u>

**Drug Requested: Repository Corticotropin Medications - Symptomatic Sarcoidosis** 

<u>PREFERRED</u>	NON-PREFERRED			
□ Purified Cortrophin <sup>™</sup> Gel	☐ HP Acthar® Gel (repository corticotropin)			
(repository corticotropin)	*Member must have tried and failed preferred Purified Cortrophin <sup>™</sup> Gel and meet all applicable PA criteria below			
MEMBER & PRESCRIBER INFORMAT	ΓΙΟΝ: Authorization may be delayed if incomplete.			
Member Name:				
Iember Sentara #: Date of Birth:				
Prescriber Name:				
Prescriber Signature: Date:				
Office Contact Name:				
Phone Number: Fax Number:				
DEA OR NPI #:				
DRUG INFORMATION: Authorization may	be delayed if incomplete.			
Drug Form/Strength/Month:				
Dosing Schedule:	Length of Therapy:			
Diagnosis:	nosis: ICD Code, if applicable:			
Diagnosis	ICD Code, ii applicable:			
<ul> <li>Adverse effects that may occur with repository effects and are similar to corticosteroids. There</li> </ul>	corticotropin are related primarily to its <u>steroidogenic</u> e may be increased susceptibility to new infection and as. Adrenal insufficiency may occur after abrupt			
Adverse effects that may occur with repository effects and are similar to corticosteroids. There increased risk of reactivation of latent infection withdrawal of the drug following prolonged the CLINICAL CRITERIA: Check below all that	corticotropin are related primarily to its <u>steroidogenic</u> e may be increased susceptibility to new infection and as. Adrenal insufficiency may occur after abrupt			
Adverse effects that may occur with repository effects and are similar to corticosteroids. There increased risk of reactivation of latent infection withdrawal of the drug following prolonged the CLINICAL CRITERIA: Check below all that each line checked, all documentation, including lab	corticotropin are related primarily to its steroidogenic e may be increased susceptibility to new infection and as. Adrenal insufficiency may occur after abrupt erapy.  t apply. All criteria must be met for approval. To support results, diagnostics, and/or chart notes, must be provided or			
Adverse effects that may occur with repository effects and are similar to corticosteroids. There increased risk of reactivation of latent infection withdrawal of the drug following prolonged the CLINICAL CRITERIA: Check below all that each line checked, all documentation, including lab request may be denied.	corticotropin are related primarily to its steroidogenic e may be increased susceptibility to new infection and as. Adrenal insufficiency may occur after abrupt erapy.  t apply. All criteria must be met for approval. To support results, diagnostics, and/or chart notes, must be provided or			

	Member <u>must</u> have tried and failed or has a contraindication to systemic corticosteroids as follows:				
	☐ Trial of dose equivalent to at least 20 mg prednisone daily for 3 months <u>MUST</u> be noted in pharmacy claims				
	OR				
	☐ For contraindication: GI BLEED has occurred within the last 30 days (must submit chart note documentation)				
	AND				
	Member must have tried and failed or has a contraindication to at least <u>one</u> (1) of the following immunomodulators (therapy tried <u>must</u> be noted in pharmacy claims):				
	□ methotrexate	□ azathioprine	□ leflunomide		
	AND				
	Member must have tried and failed or has a contraindication to at least <u>one</u> (1) TNF Inhibitor (therapy tried <u>must</u> be noted in pharmacy claims):				
	□ infliximab (Remicade®)	□ etanercept (Enbrel®)	□ adalimumab (Humira <sup>®</sup> )		
	AND				
	Documentation that <u>EITHER</u> pulmonary imaging/pulmonary function tests <u>OR</u> noncaseating granulomas showed worsening of disease while on a steroid and immunomodulator and TNF-Inhibitor (progress notes and diagnostics <u>MUST</u> be submitted):				
	<ul><li>Pulmonary imaging</li></ul>	OR □ Co	nfirmation of noncaseating granulomas		
	☐ Recent pulmonary function te	ests			
Me	Medication being provided by a Specialty Pharmacy - PropriumRx				

Not all drugs may be covered under every Plan

If a drug is non-formulary on a Plan, documentation of medical necessity will be required.

\*\*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. \*

<sup>\*</sup>Approved by Pharmacy and Therapeutics Committee: 7/21/2016 REVISED/UPDATED/REFORMATTED: 44/8/2020; 6/46/2022; 10/26/2023