## **HEALTHY U CHIP**

#### PRIOR AUTHORIZATION REQUEST FORM

### **Eosinophilic Granulomatosis with Polyangiitis (EPGA)**

Fasenra®, Nucala®

For authorization, please answer each question and fax this form PLUS chart notes back to the Healthy U CHIP Prior Authorization Department.

- For Medical Pharmacy please fax requests to: 801-213-1547
- For **Retail Pharmacy** please fax requests to: 385-425-4052

Failure to submit clinical documentation to support this request will result in a dismissal of the request.

If you have prior authorization questions, please call for Pharmacy Customer Service for assistance at 385-425-5094

	Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.								
Dat	re:	Member Name:		ID#:					
DO	В:	Gender:		Physi	sician:				
Off	ice Phone:	Office Fax:		Offic	Office Contact:				
Height/Weight:			HCPCS Code:						
pre rea	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.  Preferred/Non-Preferred  1. Preferred  2. Fasenra® (benralizumab)  2. Non-Preferred  a. Nucala® (mepolizumab)								
Product being request:									
Dosing/Frequency:									
If the request is for reauthorization, proceed to reauthorization section									
Questions			o reaut	IIOIIZat	ion section				
	Question		Yes	No	Comments/Notes				
1.	Is the request made by, or in cons rheumatologist, allergist, or immu	ns ultation with, a pulmonologist,							
1. 2.	Is the request made by, or in cons	ultation with, a pulmonologist, nologist?	Yes	No					
	Is the request made by, or in cons rheumatologist, allergist, or immu Does the member have a past me	ultation with, a pulmonologist, nologist? dical history or presence of	Yes	No	Comments/Notes				

	Palpable purpura	infiltration or eosinophil							
		rich granulomatous							
		inflammation							
5.	Has the member been on a stable co				Please provide documentation				
	4 weeks prior to initiating the requested therapy?								
6.					Please provide documentation				
	immunosuppressants used for main	tenance therapy:							
	azathioprine, methotrexate, or leflu	nomide?							
7.	Does documentation show objective	e baseline severity (e.g.			Please provide documentation				
	nighttime awakenings, daytime sym	ptoms, FEV1, etc.)?							
REAUTHORIZATION									
1.	Is the request for reauthorization of	therapy?							
2.	Does updated documentation show				Please provide documentation				
	experienced a positive clinical respo				·				
	following:								
	<ul> <li>reduction in the frequency and/o</li> </ul>	or severity of relapses							
	<ul> <li>reduction or discontinuation of c</li> </ul>								
	and/or immunosuppressants								
	<ul> <li>disease remission</li> </ul>								
		cy of ECDA-related symptoms							
<ul> <li>reduction in severity or frequency of EGPA-related symptoms</li> <li>What medications and/or treatment modalities have been tried in the past for this condition? Please document</li> </ul>									
			ie past	ior tilis	condition? Please document				
name of treatment, reason for failure, treatment dates, etc.									
Additional information:									
Physician's Signature:									

# \*\* Failure to submit clinical documentation to support this request will result in a dismissal of the request.\*\*

Policy: PHARM-CHIP-163 Origination Date: 01/23/2025 Reviewed/Revised Date: 01/29/2025 Next Review Date: 01/29/2026 Current Effective Date: 02/01/2025

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