## **AvMed**

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

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid)</u> on this request. All other information may be filled in by office staff; <u>fax to 1-305-671-0200</u>. No additional phone calls will be necessary if all information (<u>including phone and fax #s</u>) on this form is correct. <u>If the information provided is not complete, correct, or legible, the authorization process can be delayed.</u>

Drug Requested: Nucala® (mepolizumab) (Pharmacy)

{Hypereosinophilic Syndrome (HES)}

MEMBER & PRESCRIBER IN	<b>FORMATION:</b> Authorization may be delayed if incomplete.
Member Name:	
Member AvMed #:	Date of Birth:
Prescriber Name:	
Prescriber Signature:	Date:
Office Contact Name:	
Phone Number:	Fax Number:
DEA OR NPI #:	
DRUG INFORMATION: Author	rization may be delayed if incomplete.
Drug Form/Strength:	
Dosing Schedule:	Length of Therapy:
Diagnosis:	ICD Code, if applicable:
Weight:	Date:
Recommended Dosage: 300 mg/mL Sub	oQ once every 4 weeks administered as 3 separate 100-mg injections
Tezspire® and Xolair® to be experimentave NOT been established and will No	concomitant therapy with Cinqair <sup>®</sup> , Dupixent <sup>®</sup> , Fasenra <sup>®</sup> , Nucala <sup>®</sup> atal and investigational. Safety and efficacy of these combinations OT be permitted. In the event a member has an active Cinqair <sup>®</sup> , air <sup>®</sup> authorization on file, all subsequent requests for Nucala <sup>®</sup> will
	below all that apply. All criteria must be met for approval. To ration, including lab results, diagnostics, and/or chart notes, must be
Initial Authorization: 12 month	S
$\square$ Member is $\ge 12$ years of age	

(Continued on next page)

☐ Prescriber is or has consulted with an Allergist, Immunologist, Pulmonologist or Rheumatologist

	Member has a diagnosis of HES for 6 months or longer without any non-hematologic secondary cause (i.e. drug hypersensitivity, parasitic helminth infection, human immunodeficiency virus infection, non-hematologic malignancy) (submit chart notes and labs confirming diagnosis)	
	Member has FIP1L1-PDGFRα-negative disease	
	Member has had two or more episodes of HES-related flares (worsening of clinical symptoms and/or worsening of blood eosinophil counts) requiring escalation of therapy in the past 12 months (submit chart notes)	
	Member's HES-related flares occur spontaneously and did <u>NOT</u> occur within 4 weeks of a decrease in therapy	
	Member has been on a stable dose of HES therapy (such as oral corticosteroids, immunosuppressive agents and/or cytotoxic therapy) for the past 4 or more weeks (verified by pharmacy paid claims)	
	Member's blood eosinophil count is $\geq 1000$ cells/microliter while taking stable doses of HES therapy (submit labs obtained within 4 weeks of request)	
uppc	uthorization: 12 months. Check below all that apply. All criteria must be met for approval. To ort each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be ded or request may be denied.	
	Member has experienced a positive response to Nucala® therapy as determined by the prescriber (i.e. decreased number of flares, improved fatigue, reduced corticosteroid requirements, and decreased eosinophil levels) (submit chart notes)	
Medication being provided by a Specialty Pharmacy – Proprium Rx		

\*\*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. \*