SENTARA COMMUNITY PLAN (MEDICAID)

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 1-800-750-9692</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>

<u>Drug Requested</u>: Nucala® (mepolizumab) (Pharmacy) (Non-Preferred)

Chronic rhinosinusitis with nasal polyps (CRSwNP)

MEMBER & PRESCRIBER IN	FORMATION: Authorization may be delayed if incomplete.
Member Name:	
Member Sentara #:	
Prescriber Name:	
	Date:
Office Contact Name:	
Phone Number:	Fax Number:
DEA OR NPI #:	
DRUG INFORMATION: Authoriz	zation may be delayed if incomplete.
Drug Name/Form/Strength:	
Dosing Schedule:	Length of Therapy:
Diagnosis:	ICD Code, if applicable:
Weight:	Date:

Recommended Dosage:

• 100 mg/mL subcutaneously once every 4 weeks

Quantity Limit: 100 mg per 28 days

*Sentara Health Plans considers the use of concomitant therapy with Cinqair®, Dupixent®, Fasenra®, and Xolair® to be experimental and investigational. Safety and efficacy of these combinations have <u>NOT</u> been established and will <u>NOT</u> be permitted. In the event a member has an active Cinqair®, Dupixent®, Fasenra®, and/or Xolair® authorization on file, any subsequent requests for Nucala® will <u>NOT</u> be approved.

CLINICAL CRITERIA: Check below all that apply. <u>All criteria must be met for approval.</u> To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

Initial Authorization: 6 months 1. Has the member been approved for Nucala* previously through the Sentara medical department? Yes	□ D	IAGNOSIS: Chronic Rhinosinusitis with Nasal Polyps (CRSwNP)
 □ Yes □ No 2. Is the member 18 years of age or older? □ Yes □ No 3. Does the member have bilateral symptomatic sino-nasal polyposis with symptoms lasting at least 8 weeks? □ Yes □ No 4. Has the member failed at least 8 weeks of intranasal corticosteroid therapy? □ Yes □ No 5. Will therapy be used in combination with intranasal corticosteroids unless unable to tolerate or contraindicated? □ Yes □ No 6. Has the member tried and failed an adequate trial of the preferred product Xolair*? □ Yes □ No 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. 1. Has the member been assessed for toxicity? □ Yes □ No 2. Does the member have disease response as indicated by improvement in signs and symptoms compare baseline in one or more of the following: nasal/obstruction symptoms, improvement of sinus opacifications as assessed by CT-scans and/or an improvement on a disease activity scoring tool [e.g., nasal polyposis score (NPS), nasal congestion (NC) symptom severity score, sinonasal outcome test-22 (SNOT-22), etc.]? (supporting chart notes submitted) □ Yes □ No 3. Has the member had improvement in at least one of the following: reduction in nasal polyp size, reducting the need for systemic corticosteroids, improvement in quality of life, improvement in sense of smell and/or reduction of impact of comorbidities? (supporting chart notes submitted) □ Yes □ No Medication being provided by (check box below that applies): 		
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Medication being provided by (check box below that applies):	3.	
□ Physician's office OR □ Specialty Pharmacy - PropriumRx	Med	ication being provided by (check box below that applies):
		Physician's office OR Specialty Pharmacy - PropriumRx
**Use of samples to initiate therapy does not meet step edit/ preauthorization criteria. **	* *	KUsa of samples to initiate thoughy does not most stap adit/proceeds origination oritorie **

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