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® SQ (mepolizumab) (<u>Pharmacy</u>) (<u>Non-Preferred</u>) Eosinophilic Granulomatosis Polyangiitis (EGPA)*

MEMBER & PRESCRIBER INF	TORMATION: Authorization may be delayed if incomplete.						
Member Name:							
Member Sentara #:	er Sentara #: Date of Birth:						
Prescriber Name:							
Prescriber Signature:							
Office Contact Name:							
	Fax Number:						
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 (if applicable):	Date weight obtained:						

Recommended Dosage: 300 mg/mL SubQ once every 4 weeks administered as 3 separate 100-mg injections; single-dose prefilled auto-injector/single-dose prefilled syringe

*The Health Plan considers the use of concomitant therapy with Cinqair[®], Dupixent[®], Fasenra[®], Tezspire[™] 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[®], Tezspire[™] or Xolair[®] authorization on file, all subsequent requests for Nucala[®] will <u>NOT</u> be approved.

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CLINICAL CRITERIA: 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.

<u>Initi</u>	al Authorization: 6 months
1.	Has the member been approved for Nucala [®] previously through the Sentara medical department? ☐ Yes ☐ No
2.	Is the member 18 years of age or older? □ Yes □ No
3.	Does the member have a confirmed diagnosis of EGPA (aka Churg-Strauss Syndrome)? □ Yes □ No
4.	Does the member have blood eosinophils ≥ 1000 cells/ μL or $\geq 10\%$ eosinophils on white blood cell differential count?
5.	 □ Yes □ No Has the member been on stable dose of concomitant oral corticosteroid therapy for at least 4 weeks (i.e. prednisone or prednisolone at a dose of 7.5 mg/day)? □ Yes □ No
6.	Has the physician assessed baseline disease severity utilizing an objective measure/tool (e.g., Birmingham Vasculitis Activity Score [BVAS], history of asthma symptoms and/or exacerbations, duration of remission, rate of relapses)? □ Yes □ No
7.	Has the member tried and failed and adequate trial of the preferred product Fasenra®? □ Yes □ No
suppo	Ithorization: 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.
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 compared to baseline as evidenced in one or more of the following:
	• Member is in remission [defined as a Birmingham Vasculitis Activity Score (BVAS) score=0 and a prednisone/prednisolone daily dose of ≤ 7.5 mg]
	Decrease in maintenance dose of systemic corticosteroids
	Improvement in BVAS score compared to baseline
	• Improvement in asthma symptoms or asthma exacerbations
	• Improvement in duration of remission or decrease in the rate of relapses?
	□ Yes □ No

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*Eosinophilic Granulomatosis Polyangiitis (EGPA) is defined as all the following:

- History or presence of asthma
- Blood eosinophil level > 10% or an absolute count > 1000 cells/mm³
- Two or more of the following criteria:
 - o Histopathologic evidence of eosinophilic vasculitis, perivascular eosinophilic infiltration, or eosinophil rich granulomatous inflammation
 - Neuropathy
 - o Pulmonary infiltrates
 - Sinonasal abnormalities
 - Cardiomyopathy
 - Glomerulonephritis
 - Alveolar hemorrhage
 - o Palpable purpura
 - o Antineutrophil Cytoplasmic Antibody (ANCA) positivity

Medication	being	provided by	Specialty	y Pharmacy	v - Pro	priumRx
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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.