## SENTARA HEALTH PLANS

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

<u>Directions:</u> The prescribing physician must sign and clearly print name (preprinted stamps not valid) on this request. All other information may be filled in by office staff; <u>fax to 1-844-668-1550</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 information provided is not complete, correct, or legible, authorization can be delayed</u>.

<u>For Medicare Members:</u> Medicare Coverage for outpatient (Part B) drugs is outlined in the Medicare Benefit Policy Manual (Pub. 100-2), Chapter 15, §50 Drugs and Biologicals. In addition, National Coverage Determination (NCD) and Local Coverage Determinations (LCDs) may exist and compliance with these policies is required where applicable. They can be found at: <a href="https://www.cms.gov/medicare-coverage-database/overview-and-quick-search.aspx">https://www.cms.gov/medicare-coverage-database/overview-and-quick-search.aspx</a>. Additional indications may be covered at the discretion of the health plan.

<u>Drug Requested</u>: Nucala® (mepolizumab) (J2182) (Medical)

**Chronic Obstructive Pulmonary Disease (COPD)** 

MEMBER & PRESCRIBER INFORMATION: Authorization may be delayed if incomplete.				
Member Name:				
Member Sentara #:				
Prescriber Name:				
Prescriber Signature:				
Office Contact Name:				
Phone Number:	Fax Number:			
NPI #:				
DRUG INFORMATION: Authoriza	ntion 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:			
	the timeframe does not jeopardize the life or health of the member turn function and would not subject the member to severe pain.			

**Quantity Limit:** 100 mg per 28 days

\*The Health Plan considers the use of concomitant therapy with Cinqair®, Nucala®, Dupixent®, Fasenra®, and Xolair® to be experimental and investigational. Safety and efficacy of these combinations have NOT been established and will NOT 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 NOT be approved.

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<b>CLINICAL CRITERIA:</b> 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.					
	□ DIAGNOSIS: Chronic Obstructive Pulmonary Disease				
<u>Initial Authorization</u> : 12 months					
	Prescribed by or in consultation with an allergist, immunologist or pulmonologist				
	Member is 18 years of age or older				
	Member has a diagnosis of moderate to severe Chronic Obstructive Pulmonary Disease (COPD) confirmed with spirometry demonstrating ONE of the following:				
	FEV1/FVC ratio <0.7 post-bronchodilation  Rest bronchodilator FEV1 % and istad of > 20% and < 80%				
_	Post-bronchodilator FEV1 % predicted of $\geq 30\%$ and $\leq 80\%$				
	Member has experienced <u>ONE</u> of the following:  ☐ At least two (2) exacerbations treated with short-acting bronchodilators and oral corticosteroids, with or without antibiotics in the past 12 months				
	☐ At least one (1) exacerbation requiring hospitalization in the past 12 months				
	Provider must submit a member blood eosinophil count level greater than or equal to 300 cells per microliter following at least 90 days of therapy of dual or triple-maintenance therapies				
	Member is currently being treated with <u>ONE</u> of the following unless there is a contraindication or intolerance to these medications and must be compliant on therapy <u>for at least 90 consecutive days</u> within year of the request (verified by pharmacy paid claims):				
	☐ Triple therapy with a long-acting muscarinic antagonist (LAMA) (e.g., Spiriva Respimat®), long-acting beta agonist (LABA) (e.g., Advair HFA, Dulera®), and an inhaled corticosteroid (ICS) (e.g., fluticasone propionate)				
	Dual therapy with a long-acting muscarinic antagonist (LAMA) (e.g., Spiriva Respimat®) and long-acting beta agonist (LABA) (e.g., Advair HFA, Dulera®) alone if inhaled corticosteroid (ICS) is contraindicated				
	Member is requesting Nucala® (mepolizumab) as add-on maintenance therapy to dual or triple therapy (verified by pharmacy paid claims)				

**Reauthorization: 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.

		ember has experienced a sustained positive clinical response to Nucala® therapy as monstrated by at least <b>ONE</b> of the following (check all that apply):
		Increase in percent predicted Forced Expiratory Volume (FEV1) from baseline (pretreatment)
		Reduction in exacerbations (e.g., decrease oral corticosteroids) or fewer hospitalizations
		Reduction in dyspnea symptoms such as chest tightness, shortness of breath

PA Nucala (COPD) (Medical) (CORE) (continued from previous page)

_	Member is currently being treated with ONE of the following unless there is a contraindication or intolerance to these medications (verified by pharmacy paid claims):  ☐ Triple therapy with a long-acting muscarinic antagonist (LAMA) (e.g., Spiriva Respimat®), long-acting beta agonist (LABA) (e.g., Advair HFA, Dulera®), and an inhaled corticosteroid (ICS) (e.g., fluticasone propionate)  ☐ Dual therapy with a long-acting muscarinic antagonist (LAMA) (e.g., Spiriva Respimat®) and long-acting beta agonist (LABA) (e.g., Advair HFA, Dulera®) alone if inhaled corticosteroid (ICS) is
	contraindicated  Member continues to use Nucala® (mepolizumab) as add-on maintenance therapy to dual or triple therapy (verified by pharmacy paid claims)
Me	ication being provided by (check applicable box(es) below):
	Physician's office OR   Specialty Pharmacy
standa urgent	ent reviews: Practitioner should call Sentara Health Plans Pre-Authorization Department if they believe a dreview would subject the member to adverse health consequences. Sentara Health Plan's definition of s a lack of treatment that could seriously jeopardize the life or health of the member or the member's o regain maximum function.
	Use of samples to initiate therapy does not meet step edit/preauthorization criteria.**  ious therapies will be verified through pharmacy paid claims or submitted chart notes.