

Referral Status:	MRN:
<input type="checkbox"/> New referral	<input type="checkbox"/> Order change
<input type="checkbox"/> Order Renewal	
Patient preferred clinic:	

**Nucala® (mepolizumab) Pediatric (aged 6 to 11 years) Standard Plan of Treatment for Asthma**

Rev 4.29.25

**PATIENT DEMOGRAPHICS:**

Patient Name:	
Patient's Phone:	Address:
Date of Birth:	City, State, Zip:
Height in inches:	Weight: LB or KG Gender: Allergies: See list NKDA

**DIAGNOSIS: (PLEASE COMPLETE 2<sup>ND</sup> AND 3<sup>RD</sup> DIGITS TO COMPLETE ICD 10 FOR BILLING)**

<input type="checkbox"/> J45.50 - Severe persistent asthma, uncomplicated
<input type="checkbox"/> J45.52 - Severe persistent asthma with status asthmaticus
<input type="checkbox"/> J45.51 - Severe persistent asthma with (acute) exacerbation
<input type="checkbox"/> - Other:

**REQUESTED DOCUMENTATION:**

1	Insurance information
2	Most recent History & Physical
3	Full medication list
4	Tried and failed therapies
5	Blood eosinophil level (pre-treatment baseline count greater than or equal to 150 cells/mcL)

**PREVIOUS ADMINISTRATION: HAS THIS PATIENT TAKEN THIS MEDICATION BEFORE?**

IF NO:	IF YES:
PLEASE STATE REQUIRED WASHOUT FROM PREVIOUS THERAPY:	LAST INJECTION DATE:
	NEXT INJECTION DATE:
<b>IF ORDER CHANGE:</b>	
<b>Continue current order until insurance approved</b>	

**Provider Attestation for HCP administration:**

<input type="checkbox"/> Provider attestation that the patient or caregiver are not competent or are physically unable to administer the Nucala product FDA labeled for self-administration	<input type="checkbox"/> Patient has experienced severe hypersensitivity reactions (e.g., anaphylaxis, angioedema, bronchospasm, or hypotension) to Nucala within the past 6 months and requires administration and direct monitoring by a healthcare professional*
<input type="checkbox"/> Patient has a history of uncontrolled disease and ordering provider attests that in their clinical opinion, it is not advisable to try the self-administered formulation of requested drug	<input type="checkbox"/> Due to patient's weight, ordering provider attests that in their clinical opinion, it is not advisable to try the self-administered formulation of requested drug
<input type="checkbox"/> The location and circumstances for self-administration are not adequate for the potential treatment of anaphylaxis should that arise.	

\*Specific reactions: \_\_\_\_\_

**MEDICATION ORDERS:**

NOTE: Patient may be ineligible to receive Nucala® (mepolizumab) if patient has signs/symptoms of parasitic infection, is currently being treated for a parasitic infection, or is having acute bronchospasm and/or asthma attack.

**DOSE/FREQUENCY:**

Nucala® (mepolizumab) 40 mg every four (4) weeks via subcutaneous injection

**Administer as subcutaneous injection to the upper arm, thigh, or abdomen.**

**SPECIAL ORDERS:**

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**Extended post treatment monitoring: monitor patient for one (1) hour after first injection, 30 minutes after second injection, and 15 minutes after each subsequent injection.**

Refills x 12 months unless noted otherwise here:

**NURSING ORDERS:**

Provide nursing care per Palmetto Infusion Nursing Procedures and post procedure observation if indicated.

**ADVERSE REACTION & ANAPHYLAXIS ORDERS:**

Administer acute infusion and anaphylaxis medications per Palmetto Infusion standing adverse reaction orders, which can be found at our website.

**PRESCRIBER INFORMATION:**

PROVIDER NAME:	PHONE:
ADDRESS:	FAX:
CITY, STATE, ZIP:	NPI:

**PRESCRIBER SIGNATURE: (No stamp signatures)**

**DATE**

Dispense as written/Brand medically necessary	Substitution permitted