

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

## Nucala® (mepolizumab) Standard Plan of Treatment for Hypereosinophilic Syndrome (HES)

### 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)

D72.119 - Hypereosinophilic syndrome (HES), unspecified

- Other: \_\_\_\_\_

REQUESTED DOCUMENTATION:	PREVIOUS ADMINISTRATION: HAS THIS PATIENT TAKEN THIS MEDICATION BEFORE?
1 Insurance information	IF NO: _____ IF YES: _____
2 Most recent History & Physical	PLEASE STATE LAST INJECTION DATE: _____
3 Full medication list	REQUIRED WASHOUT FROM PREVIOUS THERAPY: _____ NEXT INJECTION DATE: _____
4 Tried and failed therapies	<b>IF ORDER CHANGE:</b> <input type="checkbox"/> <b>Continue current order until insurance approved</b>
5 Blood eosinophil level (pre-treatment baseline count greater than or equal to 150 cells/mcL)	

#### 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) 300 mg every four (4) weeks via subcutaneous injection for the treatment of adult and pediatric patients aged 12 years and older with hypereosinophilic syndrome (HES).

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

### SPECIAL ORDERS:

\_\_\_\_\_

**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