

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 EGPA

### PATIENT DEMOGRAPHICS:

Date of Referral:	Patient's Phone:
Patient Name:	Address:
Date of Birth:	City, State, Zip:
Height in inches:	Weight: LB or KG
Gender:	Allergies:
<input type="checkbox"/> See list	<input type="checkbox"/> NDKA

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

M30.1 - Polyarteritis with lung involvement (Eosinophilic Granulomatosis with Polyangiitis: Churg Strauss Syndrome)
- Other:

### REQUESTED DOCUMENTATION:

1	Insurance information
2	Most recent History & Physical
3	Full medication list
4	Tried and failed therapies

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

IF NO:	IF YES:
PLEASE STATE	LAST INJECTION DATE:
REQUIRED WASHOUT	NEXT INJECTION DATE:
FROM PREVIOUS	
THERAPY:	
	<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) 300 mg every four (4) weeks via subcutaneous injection

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:

### ADVERSE REACTION & ANAPHYLAXIS ORDERS:

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



### 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	