

## PATIENT INFORMATION

PATIENT NAME: \_\_\_\_\_ PHONE: \_\_\_\_\_

DATE OF BIRTH: \_\_\_\_\_ SEX: ☐ M ☐ F HEIGHT: \_\_\_\_\_ WEIGHT: \_\_\_\_\_ ☐ LBS ☐ KG

ALLERGIES: \_\_\_\_\_ PREFERRED CLINIC: \_\_\_\_\_

REFERRAL STATUS: ☐ NEW REFERRAL ☐ ORDER CHANGE ☐ ORDER RENEWAL

## DIAGNOSIS & CLINICAL DOCUMENTATION

\*PLEASE COMPLETE ICD-10 FOR SPECIFIC DIAGNOSIS

- |   |   |
|---|---|
| <input type="checkbox"/> D72. ____ Hypereosinophilic Syndrome – HES | <input type="checkbox"/> J82.83 Eosinophilic Asthma                                       |
| <input type="checkbox"/> J33. ____ Nasal Polyps                     | <input type="checkbox"/> M30.1 Polyarteritis with lung involvement (EGPA / Churg-Strauss) |
| <input type="checkbox"/> J45. ____ Persistent Asthma                |   |
| <input type="checkbox"/> ICD-10 CODE: _____                         | DESCRIPTION: _____  |

## REQUIRED DOCUMENTATION

- |  |  |   |   |
|--|--|---|---|
| <input type="checkbox"/> Insurance Information                         | <input type="checkbox"/> List of Medications | <input type="checkbox"/> Tried & Failed Therapies | <input type="checkbox"/> Most Recent History & Physical |
| <input type="checkbox"/> Pulmonary Function Tests to support diagnosis | <input type="checkbox"/> Eosinophil Levels   |   |   |

## MEDICATION ORDER

\*Please complete dose when selecting frequency

- ☐ Nucala®(mepolizumab) 100mg via subcutaneous injection every 4 weeks (Asthma & Nasal Polyps dosing)
- ☐ Nucala® (mepolizumab) 300mg via subcutaneous injection every 4 weeks (EGPA & HES dosing)
- ☐ Nucala® (mepolizumab) 40mg via subcutaneous injection every 4 weeks (age 6 to 11 dosing)
- ☒ REFILL X 12 MONTHS UNLESS OTHERWISE NOTED HERE: \_\_\_\_\_

### Provider Attestation of need for HCP administration

- |                |  |
|----------------|--|
| INITIALS _____ | Patient has experienced severe hypersensitivity reactions to Nucala within the past 6 months and requires administration and direct monitoring by a healthcare professional. |
| INITIALS _____ | Patient or caregiver are not competent or are physically unable to administer the Nucala formulation labeled for self-administration.  |
| INITIALS _____ | Patient has history of uncontrolled disease and in the clinical opinion of the ordering provider, it is not advisable to try the self-administered formulation of Nucala.    |
| INITIALS _____ | The circumstances and location for self-administration are not adequate for the potential treatment of anaphylaxis.  |
| INITIALS _____ | Patient's weight is such that in the clinical opinion of the ordering provider, it is not advisable to try the self-administered formulation of Nucala.                      |

Patient to be observed for 30 minutes following the first injection and then for 15 minutes following each subsequent injection.  
In the event of an adverse reaction occurring in the infusion clinic, utilize the Immersiv Health adverse reaction protocol.

## PRESCRIBER INFORMATION

PROVIDER NAME: \_\_\_\_\_ NPI #: \_\_\_\_\_

EMAIL: \_\_\_\_\_ PHONE: \_\_\_\_\_ FAX: \_\_\_\_\_

ADDRESS (INCLUDE CITY, STATE, ZIP): \_\_\_\_\_

SUPERVISING PHYSICIAN: \_\_\_\_\_ CONTACT NAME: \_\_\_\_\_  
(IF APPLICABLE)

SIGNATURE: \_\_\_\_\_ DATE: \_\_\_\_\_  
(NO STAMPS)

SUBSTITUTION PERMITTED

DISPENSE AS WRITTEN