

## PATIENT INFORMATION

Patient Name: \_\_\_\_\_ DOB: \_\_\_\_\_ Phone: \_\_\_\_\_ Sex: M / F Ht: \_\_\_\_\_ Wt: \_\_\_\_\_ lbs / kg  
Primary Language: \_\_\_\_\_ Allergies: \_\_\_\_\_  
Patient Preferred Location: \_\_\_\_\_

&lt;ICD 10 CODE REQUIRED&gt;

## DIAGNOSIS & CLINICAL INFORMATION

**ICD 10 Code**

- J45.50 Severe Persistent Asthma, Uncomplicated  
 J45.51 Severe Persistent Asthma, w/ Acute Exacerbation  
 J45.52 Severe Persistent Asthma, w/ Status Asthmaticus  
Other: \_\_\_\_\_

**Prescribing Information**

The patient may not be eligible to receive Cinqair if they have signs, symptoms, or are being treated for a parasitic infection or if they are having acute bronchospasm and/or an asthma attack.

**REQUIRED:** Demographics & Most Recent: H&P, clinical notes, & medication list. Supporting clinical notes to include any past tried and/or failed therapies, intolerance, outcomes, or contraindications to conventional therapy.

**LAB RESULTS:** Blood Eosinophil Level (Pre-treatment baseline count  $\geq$  to 400 cells/mcL) (Absolute Eosinophil in K/mcL x1000 = cells/mcL)

## PRESCRIPTION

**Cinqair (reslizumab)**Dose:

**IV:** Infuse 3 mg/kg in 50-100 mL of 0.9% Sodium Chloride over at least  
30 minutes via pump with a 0.2-micron filter every 4 weeks for one year

**Patient Weight:** \_\_\_\_\_ lbs or \_\_\_\_\_ kg

In the event of an adverse reaction occurring at a Medix Infusion suite, utilize the Medix Infusion adverse reaction protocol.

**Post Treatment Observations:** The patient is observed for 30 minutes following the first infusion.

**Comments:** \_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_  
\_\_\_\_\_

## PRESCRIBER INFORMATION

Prescriber Name: \_\_\_\_\_ **Signature:** \_\_\_\_\_

Date: \_\_\_\_\_ NPI #: \_\_\_\_\_ Specialty: \_\_\_\_\_

Supervising Physician: \_\_\_\_\_ (If Applicable)

Address: \_\_\_\_\_ City: \_\_\_\_\_ State: \_\_\_\_\_ Zip: \_\_\_\_\_

Contact Name: \_\_\_\_\_ Phone: \_\_\_\_\_ Fax: \_\_\_\_\_ Email: \_\_\_\_\_