

**CINQAIR (RESILUZUMAB)  
(NON-PREFERRED)  
PRIOR AUTHORIZATION FORM**  
(form effective 1/5/21)



**Keystone First**

**PERFORMRx**  
Next Generation Pharmacy Benefits

Fax to PerformRx<sup>SM</sup> at **1-215-937-5018**, or to speak to a representative call **1-800-588-6767**.

**PRIOR AUTHORIZATION REQUEST INFORMATION**

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request	total # pages:	Name of office contact:
Contact's phone number:	LTC facility contact/phone:	

**PATIENT INFORMATION**

Patient name:	Patient ID #:	DOB:
Street address:	Apt. #:	City/state/zip:

**PRESCRIBER INFORMATION**

Prescriber name:	Specialty:	
State license #:	NPI:	MA Provider ID #:
Street address:	Suite #:	City/state/zip:
Phone:	Fax:	

**CLINICAL INFORMATION**

**Medication requested:**  Cinqair 100 mg/10 ml vial     Cinqair \_\_\_\_\_

**Dose/directions:** \_\_\_\_\_

<b>Quantity requested:</b> # _____ vials (100 mg/10 ml vial)	<b>Duration requested:</b> _____ months	<b>Weight:</b> _____ lbs / kg
<b>Diagnosis:</b> _____		<b>Dx code (required):</b> _____

**PHARMACY INFORMATION (Prescriber to identify the pharmacy that is to dispense the medication):**

Deliver to:  Patient's Home     Physician's Office     Patient's Preferred Pharmacy Name:

Pharmacy Phone #: \_\_\_\_\_ Pharmacy Fax #: \_\_\_\_\_

I acknowledge that the patient agrees with the pharmacy chosen for delivery of this medication.

**INITIAL REQUESTS**

- Is Cinqair being prescribed by or in consultation with a specialist, such as a pulmonologist?  Yes     No    *Provide specialty.* \_\_\_\_\_
- Is the patient being treated for a diagnosis of asthma that is severe despite use of tolerated asthma controller medications?  Yes     No    *Submit documentation.*
- Does the patient have asthma of an eosinophilic phenotype with an absolute blood eosinophil count  $\geq 400$ /microliter?  
 Yes     No    Eosinophil count: \_\_\_\_\_ Date of result: \_\_\_\_\_
- Is the patient currently receiving and **will continue to receive** optimally titrated doses, or have a contraindication or intolerance to, any of the following?  
 inhaled glucocorticoid     leukotriene modifier     long-acting beta-agonist (LABA)     other (e.g., tiotropium, theophylline): \_\_\_\_\_  
 Yes – List medications tried: \_\_\_\_\_  No
- Does the patient have a history of trial and failure of, or contraindication or intolerance to, the preferred Monoclonal Antibodies, Anti-IL, Anti-IgE, Nucala, Xolair, and Fasenna?  
 Yes – List medications tried: \_\_\_\_\_  No
- Has the patient been using Cinqair in the past 90 days?  Yes     No    *Submit documentation.*
- Will the patient be monitored and/or treated for helminth infection as recommended in package labeling?  Yes     No

**RENEWAL REQUESTS**

- Has the patient experienced measurable evidence of improvement in asthma severity?  Yes     No    *Submit documentation of patient's response to therapy.*
- Will the patient continue to use optimally titrated doses any of the following?  
 inhaled glucocorticoid     leukotriene modifier     long-acting beta-agonist (LABA)     other (e.g., tiotropium, theophylline): \_\_\_\_\_  
 Yes     No    *Submit medical record documentation of patient's medication regimen to be used with Cinqair.*
- Does the patient have a contraindication or intolerance to optimally titrated doses of any of the medications in question 2?  
 inhaled glucocorticoid     leukotriene modifier     long-acting beta-agonist (LABA)     other (e.g., tiotropium, theophylline): \_\_\_\_\_  
 Yes     No    *Submit medical record documentation of contraindications/intolerances.*
- Will the patient be monitored and/or treated for helminth infection as recommended in package labeling?  Yes     No

**PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION**

Prescriber signature:	Date:
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