

## **MEDICARE FORM**

## Cinqair® (reslizumab) Medication Precertification Request

Page 1 of 2

(All fields must be completed and legible for precertification review.)

For New Jersey HMO D-SNP: FAX: 1-833-322-0034 PHONE: 1-844-362-0934 For other lines of business:

Please use other form

Note: Cinqair is non-preferred. The preferred products are Nucala

and Xolair.

Please indicate: ☐ Start of treatment: Start date _ ☐ Continuation of therapy: Date of		<u> </u>		and Adian.	
Precertification Requested By:		Phone:		Fax:	
A. PATIENT INFORMATION					
First Name:	Las	t Name:			
Address:	City	:		State:	ZIP:
Home Phone: Work	Phone:	C	Cell Phone:		<u>.</u>
DOB: Allergies:		E	mail:		
Current Weight: lbs or kgs	Height:	inches or	cms		
B. INSURANCE INFORMATION					
Aetna Member ID #:	Does patient have other coverage?				
Group #:	If yes, provide ID#: Carrier Name:				
Insured:	Insured:				
Medicare: ☐ Yes ☐ No If yes, provide ID #:	Med	licaid: Yes No	o If yes, pro	vide ID #:	
C. PRESCRIBER INFORMATION					
First Name:	Last Name:		(Check On	e): 🔲 M.D.	□ D.O. □ N.P. □ P.A.
Address:		City:		State:	ZIP:
Phone: Fax:	St Lic #:	NPI #:	DEA #:		UPIN:
Provider Email:	Office Contact Name:		Phone:		
Specialty (Check one): Pulmonologist Allergist Other:					
D. DISPENSING PROVIDER/ADMINISTRATION INFORMA	ATION				
Place of Administration:  Self-administered Physician's Office Outpatient Infusion Center Phone: Center Name: Home Infusion Center Phone: Agency Name:		Dispensing Provide  ☐ Physician's Office ☐ Specialty Pharm Name: Address:	ce [	Retail Pha Other	rmacy
Address:		City:	;	State:	ZIP:
Address: State:	7IP·	Phone:		Fax:	
Phone: Fax:		TIN:		PIN:	
TIN: PIN:		NPI:			
NPI:					
E. PRODUCT INFORMATION		_			
Request is for: Cinqair (reslizumab) Dose:		Frequency:			
F. DIAGNOSIS INFORMATION – Please indicate primary I		other where applicable.			
•	dary ICD Code:		Other ICD C		
G. CLINICAL INFORMATION – Required clinical information	on must be completed in it	s <u>entirety</u> for all precerti	fication reques	sts.	
For All Requests (clinical documentation required):  Note: Cinqair is non-preferred. The preferred products are Nucala, and Xolair.  Yes No Has the patient had prior therapy with Cinqair within the last 365 days?  Yes No Has the patient had a trial and failure, intolerance, or contraindication to any of the following? (select all that apply)  Nucala (mepolizumab) Xolair (omalizumab)  Please explain if there are any other medical reason(s) that the patient cannot use any of the following preferred products when indicated for the patient's diagnosis? (select all that apply)  Nucala (mepolizumab) Xolair (omalizumab)					

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB				
•	o <b>ntinued)</b> – Required clinical information	n must be completed in its <u>entiret</u>	v for all precertification requests.				
Yes No Is this infusion request in an outpatient hospital setting?							
Yes No Has the patient experienced an adverse event with the requested product that has not responded to conventional							
interventions (e.g., acetaminophen, steroids, diphenhydramine, fluids, other pre-medications or slowing of infusion rate) or a severe adverse event (anaphylaxis, anaphylactoid reactions, myocardial infarction, thromboembolism, or seizures) during or							
immediately after an infusion?							
Yes No Does the patient have severe venous access issues that require the use of special interventions only available in the							
outpatient hospital setting?							
Yes No Does the patient have significant behavioral issues and/or physical or cognitive impairment that would impact the safety of the							
infusion therapy AND the patient does not have access to a caregiver?  Please provide a description of the behavioral issue or impairment:							
☐ Yes ☐ No Is the patient medically unstable which may include respiratory, cardiovascular, or renal conditions that may limit the member's							
ability to tolerate a large volume or load or predispose the member to a severe adverse event that cannot be managed in an							
alternate setting without appropriate medical personnel and equipment?							
Please provide a description of the condition:   Cardiovascular:							
	Respiratory:						
	☐ Renal:						
		Other:					
Yes No Does the patient have a documented diagnosis of asthma?							
Yes No Will the patient receive Cinqair as monotherapy (i.e., without any other asthma medications such as inhaled corticosteroids)?							
Yes No Will the patient be taking Cinqair concomitantly with other biologics indicated for asthma (e.g., Dupixent, Fasenra, Nucala, Xolair)?							
For Initial Requests:							
Please indicate the patient's baseline (e.g., before significant oral steroid use) blood eosinophil count in cells per microliter:							
Please indicate the preferred alternatives for asthma that have been ineffective, not tolerated, or are contraindicated:   Fasenra   Nucala   Xolair  Yes   No Is the patient dependent on systemic corticosteroids?							
☐ Yes ☐ No Does the patient dependent on systemic control (e.g., hospitalization or emergency medical care visit within the past year) despite							
current treatment with both of the following medications: inhaled corticosteroid and additional controller (long acting beta-2 agonist,							
leukotriene modifier, or sustained-release theophylline) at optimized doses?							
For Continuation Requests:							
	tly receiving Cinqair through samples or a rage under the provisions of the pharmacy		orogram? (Sampling of Cinqair does				
Yes No Has asthma control improved on Cinqair treatment as demonstrated by a reduction in the frequency and/or severity of symptoms and exacerbations?							
H. ACKNOWLEDGEMENT							
Request Completed By (Signature Required): Date: //							
Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent							

insurance act, which is a crime and subjects such person to criminal and civil penalties.

The plan may request additional information or clarification, if needed, to evaluate requests.