

## Cinqair® (reslizumab) Medication Precertification Request

Page 1 of 2

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

**Aetna Precertification Notification** 

**Phone:** 1-866-752-7021 **FAX:** 1-888-267-3277

For Medicare Advantage Part B: Please Use Medicare Request Form

<ul><li>☐ Start of treatment: Start date</li><li>☐ Continuation of therapy: Date</li></ul>		/ /					
equested By:			e:	_ Fax:			
MATION							
	I	Last Name:					
	(	City:		State:	ZIP:		
We		•	Cell Phone:	1			
Allergies:			Email:				
	Height:	inches		<u> </u>			
#:	Does patient have c	other coverage?	☐ Yes ☐ No				
		=					
			_				
☐ No If yes, provide ID #:			☐ No If yes, pro	ovide ID #:			
IFORMATION							
	Last Name:		(Check On	ne): [] M.D.	☐ D.O. ☐ N.P. ☐ P.A.		
		City:		State:	ZIP:		
Fax:	St Lic #:	NPI #:	DEA #:		UPIN:		
	Office Contact Nam	ie:	<b>'</b>	Phon	e:		
ne):  Pulmonologist  Aller	aist \( \square\) Other:			l I			
·							
ed  Physician's Office sion Center Phone: me: Center Phone: ame:		☐ Physician's ☐ Specialty F ☐ Name: ☐ Address: ☐ Phone:	s Office [	Retail Pha Other  Fax:	armacy		
ıqair (reslizumab) Dose:		Frequency: _	<u></u>				
<b>DRMATION</b> – Please indicate primar	ry ICD Code and specify	any other where appl	licable.				
	<u>-</u>						
For All Requests (clinical documentation required):  Yes No Is this infusion request in an outpatient hospital setting?  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?  Does the patient have severe venous access issues that require the use of special interventions only available in the outpatient hospital setting?  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:  St 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:  Respiratory:  Respiratory:  Respiratory:							
	Continuation of therapy: Date equested By:  MATION  Allergies:	Continuation of therapy: Date of last treatment	Continuation of therapy: Date of last treatment	Continuation of therapy: Date of last treatment	Continuation of therapy: Date of last treatment		

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Patient First Na	ame	Patient Last Name	Patient Phone	Patient DOB					
G. CLINICAL	G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.								
☐ Yes ☐ No Is the medication prescribed by or in consultation with an allergist, immunologist, or pulmonologist? ☐ Yes ☐ No ☐ Does the patient have a documented diagnosis of asthma? ☐ Yes ☐ No Will the patient continue to use maintenance asthma treatments (i.e., inhaled corticosteroids, additional controller) in combination with the requested medication?									
Yes No	☐ Yes ☐ No Will the patient be taking the requested medication concomitantly with other biologics indicated for asthma (e.g., Dupixent, Fasenra, Nucala, Tezspire, 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 Does the patient have uncontrolled asthma as demonstrated by experiencing one or more asthma exacerbation resulting in hospitalization or emergency medical care visit within the past year?								
		<ul> <li>Does the patient have uncontrolled asth symptoms or reliever use, activity limited</li> </ul>	d by asthma, night waking due to	asthma) within the past year?					
Yes No Does the patient have inadequate asthma control despite current treatment with a high dose inhaled corticosteroid and additional controller (long acting beta2-agonist, long acting muscarinic antagonist, leukotriene modifier, or sustained release theophylline) at optimized doses?									
☐ Yes ☐ No	Yes No Is the patient dependent on systemic corticosteroids?								
For Continuation Requests:									
☐ Yes ☐ No	No Is the patient currently receiving the requested medication through samples or a manufacturer's patient assistance program?								
	□ No Has asthma control improved on the requested medication treatment as demonstrated by a reduction in the frequency and/or severity of symptoms and exacerbations?								
Yes No	Has asthma control improved odose?	on the requested medication treatment as	demonstrated by a reduction in t	he daily maintenance oral corticosteroid					
H. ACKNOWL	EDGEMENT.								
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.