

Fasenra® (benralizumab) Injectable Medication Precertification Request

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(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

Please indicate: Start of treatment: Start date		1 1			
Continuation of therapy: Date	e or iast treatment	<i>I I</i>	-		
Precertification Requested By: A. PATIENT INFORMATION		Phone:		ax:	
First Name:	La	st Name:			
Address:	Cit		State:	ZIP:	
Home Phone: Wo	rk Phone:	,	Cell Phone:		
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 #:		D#:Carrier Name:			
Insured:	_ Insured:				
Medicare: ☐ Yes ☐ No If yes, provide ID #:	Me	edicaid: 🗌 Yes 🔲 l	No If yes, provide ID	#:	
C. PRESCRIBER INFORMATION					
First Name:	Last Name:			M.D.	
Address:		City:	State:		
Phone: Fax:	St Lic #:	NPI #:	DEA #:	UPIN:	
Provider E-mail:	Office Contact Name:		Ph	none:	
Specialty (Check one):	gist 🗌 Other:				
D. DISPENSING PROVIDER/ADMINISTRATION INFOR	MATION				
Place of Administration: Self-administered Physician's Office Outpatient Infusion Center Phone: Center Name: Home Infusion Center Phone:		Dispensing Provider/Pharmacy: Patient Selected choice ☐ Physician's Office ☐ Retail Pharmacy ☐ Specialty Pharmacy ☐ Other:			
Agency Name:		Phone: Fax:			
Administration code(s) (CPT):				NIA I.	
Address:		TIN:	P	PIN:	
E. PRODUCT INFORMATION Request is for: Fasenra (benralizumab) Dose:		Frequency:			
F. DIAGNOSIS INFORMATION – Please indicate primary			e		
Primary ICD Code: Seco			Other ICD Code:		
G. CLINICAL INFORMATION – Required clinical informa					
For All Requests (clinical documentation required):	'				
severe adverse event (immediately after an in Yes No Does the patient have s infusion therapy AND tl Please provide a descr Yes No Is the patient medically ability to tolerate a larg alternate setting withou	nced an adverse event wit taminophen, steroids, diph anaphylaxis, anaphylactoi fusion? significant behavioral issue ne patient does not have a iption of the behavioral iss unstable which may include e volume or load or predis tt appropriate medical pers	enhydramine, fluids, oth d reactions, myocardial es and/or physical or coc ccess to a caregiver? ue or impairment: de respiratory, cardiovas pose the member to a s connel and equipment?	her pre-medications or sinfarction, thromboembers gnitive impairment that values accular, or renal conditions severe adverse event that	slowing of infusion rate) or a colism, or seizures) during or would impact the safety of the	
)		Respiratory:			
		Renal:			
	L	Journel.			

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB			
G. CLINICAL INFORMATION (continued	<i>)</i> – Required clinical information must b	pe completed in its entirety fo	r 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 Will the patient receive the requested medication concomitantly with other biologics indicated for asthma (e.g., Cinqair, Dupixent, Nucala, Tezspire, Xolair)?						
For Initiation Requests (clinical documentation required): Please indicate the patient's baseline (e.g., before significant oral steroid use) blood eosinophil count in cells per microliter: Yes No Does the patient have uncontrolled asthma as demonstrated by experiencing two or more asthma exacerbations requiring oral or injectable corticosteroid treatment within the past year? Yes No Does the patient have uncontrolled asthma as demonstrated by experiencing one or more asthma exacerbations resulting in hospitalization or emergency medical care visit within the past year? Yes No Does the patient have uncontrolled asthma as demonstrated by experiencing poor symptom control (frequent symptoms or reliever use, activity limited by asthma, night walking due to asthma) within the past year? Yes No Does the patient have inadequate asthma control despite current treatment with an inhaled corticosteroid and additional controller (long acting beta2-agonist, long-acting muscarinic antagonist, leukotriene modifier, or sustained release theophylline) at optimized doses?						
For Continuation Requests (clinical documentation required):						
☐ Yes No Is this continuation request a result of the patient receiving samples or a manufacturer's patient assistance program? ☐ Yes ☐ 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 on the requested medication treatment as demonstrated by a reduction in the daily maintenance of oral corticosteroid dose?						
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.