

Paetna ADAKVEO (crizanlizumab) Medication Precertification Request

(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:	se indicate: Start of treatment: Start date/						
	☐ Continua	ition of therapy, Date o	f last treatment/	1			
Precertification F	Requested By	y:		Phone:		Fax: _	
A. PATIENT INFO	ORMATION						
First Name:			Last Name:			DOB:	
Address:				City:		State:	ZIP:
Home Phone:		Work Phone:		Cell Phone:		Email:	
Patient Current We	eight: lb	os or kgs Patie	nt Height: inches	or cms A	llergies:		
B. INSURANCE I	NFORMATION	N					
Aetna Member ID #:			Does patient have other coverage? ☐ Yes ☐ No				
Group #:			If yes, provide ID#: Carrier Name:		arrier Name: _		
Insured:			Insured:				
Medicare: Yes		•	Medi	icaid: 🗌 Yes 🔲 I	No If yes, pro	vide ID #:	
C. PRESCRIBER	INFORMATIO	ON			(0)		
First Name:			Last Name:	Τ	(Check C	1	☐ D.O. ☐ N.P. ☐ P.A.
Address:				City:		State:	ZIP:
Phone:	Fa	ax:	St Lic #:	NPI #:	DEA #:	T	UPIN:
Provider Email:			Office Contact Name:			Phone:	
Specialty (Check	one): 🗌 Hei	matologist					
Place of Administration: Self-administered Physician's Office Outpatient Infusion Center Phone: Center Name: Home Infusion Center Phone: Agency Name: Administration code(s) (CPT): Address: PRODUCT INFORMATION				☐ Physician's C☐ Specialty Phate Name:Address:Phone:	Pensing Provider/Pharmacy: Patient Selected choice Physician's Office Retail Pharmacy Specialty Pharmacy Other me: dress: pne: Fax: PIN:		
Request is for: A	ADAKVEO (cr	izanlizumab) Dose:		_ Frequency: _			
F. DIAGNOSIS IN	IFORMATION	- Please indicate prima	ry ICD code and specify	any other where ap	oplicable.		
=			Secondary ICD Code				
G. CLINICAL INFORMATION - Required clinical information must be completed in its entirety for all precertification requests. 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? 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: 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: Cardiopulmonary: Respiratory:							
				Renal:			
•				Other:			

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Patient First Name	Patient Last Name	Patient Phone	Patient DOB					
G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.								
 Yes No Does the patient have a documented diagnosis of sickle cell disease? Yes No Is the requested medication being requested for use in reducing the frequency of vasoocclusive crises (VOCs)? Yes No Is the requested medication being prescribed by or in consultation with a hematologist or specialist in sickle cell disease? 								
For Initial Requests (clinical documentation required):								
Please indicate the sickle cell genotype:								
 Homozygous hemoglobin S (HbSS) or ☐ Sickle beta0-thalassemia (HbSbeta0) Yes ☐ No Has the patient experienced, at any time in the past, an inadequate response or intolerance to a trial of hydroxyurea? Yes ☐ No Does the patient have a contraindication to hydroxyurea? Yes ☐ No Will the patient be using the requested medication with concurrent hydroxyurea therapy? Please specify: ☐ Patient had an inadequate response ☐ Patient had an intolerance ☐ Sickle hemoglobin C (HbSC) ☐ Sickle beta+-thalassemia (HbSbeta+) 								
Other/Unknown:								
 ☐ Yes ☐ No Has the patient experienced a vasoocclusive crisis (VOC) in the past 12 months? For Continuation Requests (clinical documentation required): ☐ Yes ☐ No Has the patient experienced a reduction in the frequency of vasoocclusive crises, or has the patient maintained a reduction in the frequency of vasoocclusive crises, since initiating therapy with the requested medication? 								
H. ACKNOWLEDGEMENT								
Request Completed By (Signature	Required):		Date:/					
any insurance company by providing	quest for authorization of coverage of materially false information or concea ubjects such person to criminal and civ	als material information for the purpose						

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