

Tezspire® (tezepelumab-ekko) Medication Precertification Request

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

Aetna Precertification Notification Phone: <u>1-866-752-7021</u> (TTY: <u>711</u>)

FAX: <u>1-888-267-3277</u>

For Medicare Advantage Part B:
Please Use Medicare Request Form

Please indicate: Start of treatme		/ / f last treatment	1	<u> </u>		case Goo Moal	iodio rioquoci i omi		
Precertification Requested By:				Phone:		Fax:			
A. PATIENT INFORMATION									
First Name:		Last Name:				DOB:			
Address:			City	/:		State:	ZIP:		
Home Phone:	Work Phone:		Cel	I Phone:		Email:			
Patient Current Weight: lbs or _	kgs Patien	t Height:inches	or	cms Allergie	s:				
B. INSURANCE INFORMATION									
Aetna Member ID #:	Does patient have other coverage? ☐ Yes ☐ No								
Group #:		If yes, provide ID#:Carrier Name:							
Insured:			Insured:						
Medicare: ☐ Yes ☐ No If yes, prov	vide ID #:	Me	edic	aid: ☐ Yes ☐ No	If yes, provi	ide ID #:	_		
C. PRESCRIBER INFORMATION									
First Name:		Last Name:			(Check One	Î	D.O. N.P. P.A.		
Address:		-		City:		State:	ZIP:		
Phone: Fax:		St Lic #:		NPI #:	DEA #:	1	UPIN:		
Provider Email:		Office Contact Name:) :			Phone:			
Specialty (Check one): Pulmonole	ogist								
D. DISPENSING PROVIDER/ADMINI	STRATION INFO	RMATION							
Place of Administration:			_	Dispensing Provide	r/Pharmacy	y: Patient Sele	ected choice		
☐ Self-administered ☐ Phys	sician's Office	☐ Physician's Office			e [☐ Retail Pharmacy			
☐ Outpatient Infusion Center F	Phone:	Specialty Pharmacy			асу [☐ Other			
Center Name:		Name:							
☐ Home Infusion Center F	Phone:		_						
Agency Name:									
Administration code(s) (CPT):									
Address:		TIN: PIN:							
E. PRODUCT INFORMATION									
Request is for: Tezspire (tezepeluma	ab-ekko) Dose: _			Frequency:					
F. DIAGNOSIS INFORMATION - Plea	ase indicate primar	y ICD code and specify	fy any	y other where applica	ble.				
Primary ICD Code:		= =	_		 '				
G. CLINICAL INFORMATION - Requi	red clinical informa	ation must be complete	ed in	its entirety for all pred	certification	requests.			
For All Requests (clinical documentation									
Yes No Is this infusion request in			41. 41		. 4 1 4				
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, or other pre-medications) or a severe adverse event									
		oid reactions, myocardia	al infa	arction, thromboembolis	sm, or seizur	res) during or im	nmediately after		
	nistration? the patient have sig	mificant hehavioral issue	ec an	d/or physical or cogniti	ve imnairme	nt that would im	nact the safety of the		
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									
patient's ability to tolerate a large volume or load or predispose the patient 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:									
		Res	espiratory:						
		J Ker J ∩th	enal:ther:						
			_ Our	GI					
☐ Yes ☐ No Does the patient have a	diagnosis of asthm	a?							
☐ Yes ☐ No Will the requested drug I	be used concomitar	າtly with any other biolog	gic (e	g., Adbry, Humira, Du	pixent), or ta	rgeted synthetic	c drug (e.g., Rinvoq,		
Olumiant Otezla Xelian	iz) for the same indi	ication?							



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Patient First Na	nme	Patient Last Name	Patient Phone	Patient DOB				
G. CLINICAL	INFORMATION (continue	d) – Required clinical informati	on must be completed in its <u>entirety</u> for	r all precertification requests.				
			with an allergist/immunologist or pulmono					
For Initiation Requests (clinical documentation required):								
Yes No Has the patient previously received another biologic drug (e.g., Dupixent, Nucala) indicated for asthma?								
Yes No Does the patient have uncontrolled asthma as demonstrated by experiencing two or more asthma exacerbations requiring oral o injectable corticosteroid treatment within the past year?								
	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?							
	L	control (fre	eatient have uncontrolled asthma as dem equent symptoms or reliever use, activity ithin the past year?	onstrated by experiencing poor symptom limited by asthma, night waking due to				
	Yes No Prior to requesting the requested medication, did the patient have inadequate asthma control despite current treatment with							
	both of the	e following medications at optimiz	zed doses: a high dose inhaled corticoste	eroid and additional controller				
	(i.e., long-	acting beta2-agonist, long-acting	muscarinic antagonist, leukotriene modi	fier, or sustained-release theophylline)?				
	☐ Yes ☐ No Will the patient continue to use maintenance asthma treatments (e.g., inhaled corticosteroid, additional controller) in combination with the requested medication?							
For Continuation	on Requests (clinical docu	mentation required):						
Yes No	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 oral corticosteroid dose?							
Yes No Will the patient continue to use maintenance asthma treatments (e.g., inhaled corticosteroid, additional controller) in combination with the requested medication?								
H. ACKNOWL	EDGEMENT							
Request Com	pleted By <i>(Signature R</i> eq	juired):		Date:/				
any insurance	company by providing mat		ceals material information for the purp	th the intent to injure, defraud or deceive pose of misleading, commits a fraudulent				

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