



Fax completed form to: (855) 840-1678
 If this is an URGENT request, please call (800) 882-4462
 (800.88.CIGNA)

Tezspire (tezepelumab)

PHYSICIAN INFORMATION			PATIENT INFORMATION		
* Physician Name:			*Due to privacy regulations we will not be able to respond via fax with the outcome of our review unless all asterisked (*) items on this form are completed.*		
Specialty:	* DEA, NPI or TIN:				
Office Contact Person:			* Patient Name:		
Office Phone:			* Cigna ID:		* Date of Birth:
Office Fax:			* Patient Street Address:		
Office Street Address:			City:	State:	Zip:
City:	State:	Zip:	Patient Phone:		
Urgency: <input type="checkbox"/> Standard <input type="checkbox"/> Urgent (In checking this box, I attest to the fact that applying the standard review time frame may seriously jeopardize the customer's life, health, or ability to regain maximum function)					
Medication requested: <input type="checkbox"/> Tezspire 210 mg/1.91 mL (110 mg/mL) syringe <input type="checkbox"/> Tezspire 210 mg/1.91 mL Pen <input type="checkbox"/> other (please specify):					
ICD10:					
Dose		Quantity:		Frequency of therapy:	
Duration of therapy:					
Is this initial therapy, is the patient restarting therapy, or is the patient currently receiving the requested medication? <input type="checkbox"/> Initial therapy <input type="checkbox"/> Currently receiving the requested medication for less than 6 months <input type="checkbox"/> Currently receiving the requested medication and has been established on it for 6 or more months <input type="checkbox"/> Restarting therapy					
(if currently receiving) Will the patient continue to receive therapy with one inhaled corticosteroid OR one inhaled corticosteroid-containing combination inhaler? <input type="checkbox"/> Yes <input type="checkbox"/> No					
(if currently receiving) Has the patient responded to therapy as determined by the prescriber (Examples of a response to therapy are decreased asthma exacerbations; decreased asthma symptoms; decreased hospitalizations, emergency department, urgent care, or medical clinic visits due to asthma; improved lung function parameters; and/or a decreased requirement for oral corticosteroid therapy)? <input type="checkbox"/> Yes <input type="checkbox"/> No					
Where will this medication be obtained? <input type="checkbox"/> Accredo Specialty Pharmacy** <input type="checkbox"/> Hospital Outpatient <input type="checkbox"/> Retail pharmacy <input type="checkbox"/> Other (please specify): <input type="checkbox"/> Home Health / Home Infusion vendor <input type="checkbox"/> Physician's office stock (billing on a medical claim form) <i>**Cigna's nationally preferred specialty pharmacy</i> 					
<i>**Medication orders can be placed with Accredo via E-prescribe - Accredo (1620 Century Center Pkwy, Memphis, TN 38134-8822 NCPDP 4436920), Fax 888.302.1028, or Verbal 866.759.1557</i>					
Facility and/or doctor dispensing and administering medication: Facility Name: State: Tax ID#: Address (City, State, Zip Code):					

Where will this drug be administered?

- Patient's Home
 Physician's Office
 Hospital Outpatient
 Other (please specify):

NOTE: Per some Cigna plans, infusion of medication MUST occur in the least intensive medically appropriate setting.

Is this patient a candidate for re-direction to an alternate setting (such as alternate infusion site, physician's office, home) with assistance of a Specialty Care Options Case Manager? Yes No (provide medical necessity rationale):

What is your patient's diagnosis?

- Asthma
 Atopic Dermatitis
 Chronic Obstructive Pulmonary Disease (COPD)
 Chronic Rhinosinusitis with Nasal Polyposis (CRSwNP)
 Chronic Spontaneous Urticaria
 other (please specify):

Clinical Information:

Is the requested medication being prescribed by (or in consultation with) an allergist, immunologist, or a pulmonologist? Yes No

(if asthma) Did/Does the patient have a forced expiratory volume in 1 second (FEV1) of less than 80% predicted? Note: The reduced FEV1 should not be due to smoking-related chronic obstructive pulmonary disease Yes No

(if yes) Did/Does the patient have an FEV1/forced vital capacity (FVC) of less than 0.80? Yes No

(if asthma) Did/Does the patient have an increase of greater than 12% AND greater than 200 mL in FEV1 following the administration of a standard dose of a short-acting bronchodilator? Yes No

(if no) Did/Does your patient have an increase of greater than 12% AND greater than 200 mL in FEV1 between prescriber visits? Yes No

(if no) Did/Does the patient have an increase of greater than 12% AND greater than 200 mL in FEV1 from baseline to after at least 4 weeks of asthma treatment? Yes No

(if no) Did/Does the patient have a positive exercise challenge test? Yes No

(if no) Did/Does the patient have a positive bronchial challenge test? Yes No

(if asthma) Has the patient received at least 3 consecutive months of therapy with a medium- or high-dosed inhaled corticosteroid? Yes No

(if yes) During the time the patient received the medium- or high-dosed inhaled corticosteroid, did the patient also receive at least 3 consecutive months of therapy with at least one additional asthma controller or asthma maintenance medication? Examples of additional asthma controller or asthma maintenance medications are inhaled long-acting beta2-agonists, inhaled long-acting muscarinic antagonists, and monoclonal antibody therapies for asthma (for example, Tezspire, Cinqair [reslizumab intravenous infusion], Fasentra [benralizumab subcutaneous injection], Nucala [mepolizumab subcutaneous injection], Dupixent [dupilumab subcutaneous injection], Xolair [omalizumab subcutaneous injection]). Use of a combination inhaler containing both a medium- or high-dose inhaled corticosteroid and additional asthma controller/maintenance medication(s) would fulfill the requirement for both. Yes No

(if asthma) At baseline, did the patient experience two or more asthma exacerbations requiring treatment with systemic corticosteroids in the previous year? Note: Baseline is defined as prior to receiving the requested medication or another monoclonal antibody therapy for asthma (examples include Cinqair, Dupixent, Fasentra, Nucala, Tezspire, and Xolair). Yes No

(if no) At baseline, did the patient experience one or more asthma exacerbation(s) requiring hospitalization, an emergency department visit, or an urgent care visit in the previous year? Yes No

(if no) At baseline, did/does the patient have asthma that worsens upon tapering of oral (systemic) corticosteroid therapy? Yes No

(if asthma) Will the patient use the requested medication with other Monoclonal Antibodies? Yes No

(if yes) Please provide the clinical rationale for concurrent use of these drugs.

Additional Pertinent Information: *Please provide any additional pertinent clinical information, including: if the patient is currently on the requested drug (with dates of use) and how they have been receiving it (for example: samples, out of pocket).*

Attestation: I attest the information provided is true and accurate to the best of my knowledge. I understand that the Health Plan or insurer its designees may perform a routine audit and request the medical information necessary to verify the accuracy of the information reported on this form.

Prescriber Signature: _____ **Date:** _____

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