

## Pharmacy Request for Prior Approval - Tezspire

Beneficiary Information			
1. Beneficiary Last Name:	2. First Name:		
3. Beneficiary ID #:	4. Beneficiary Date of Birth:	5. Ben	eficiary Gender:
Prescriber Information			
6. Prescriber Name:		NPI #:	
Mailing address:	City:	State:	 ZIP:
7. Requester Contact Information:			
Name:	Phone #:	Fax #:	
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8. Drug Name:	9. Strength:	10. Quantity Per 30	Days:
11. Length of Therapy:up to 30 days	60 days90 days120 days _	180 days365 days _	Other:
Clinical Information			
Initial Approval **Initial approval can be for	up to 6 months**		
1. Is the beneficiary 12 years of age or older? Yes No			
2. Does the beneficiary have a diagnosis of severe asthma with evidence of severe disease? Yes No			
3. Does the beneficiary have at least 1 of the following? Yes No Please indicate which one:			
a. Symptoms throughout the day			
b. Nighttime awakenings, often 7x/week			
c. SABA use for symptom control occurring several times per day			
d. Extremely limited normal activities			
e. Lung function (percent predicted FEV1) < 60%			
f. Exacerbations requiring oral systemic corticosteroids generally more frequent and intense relative to moderate asthma			
4. Is Tezspire being used for add-on maintenance treatment for a beneficiary who regularly received BOTH of the following? YesNo			
a. Medium- to high-dose inhaled corticosteroids			
b. An additional controller medication (e.g., long-acting beta-agonist, leukotriene modifiers)			
5. Has the beneficiary had, in the previous year, ≥ 2 exacerbations requiring oral or injectable corticosteroid treatment (in addition to the regular maintenance thorsely defined above) OP one exacerbation regularing in beginning. Yes			
maintenance therapy defined above) <b>OR</b> one exacerbation resulting in hospitalization? Yes No 6. Is there a baseline measurement of ≥ 1 of the following for assessment of clinical status? Yes No			
Please indicate which one(s):			
a. Use of systemic corticosteroids			
b. Use of inhaled corticosteroids			
c. Number of hospitalizations, ER visits, or unscheduled visits to healthcare provider due to condition			
d. FEV1			
7. Will the beneficiary use Tezspire for the relief of acute bronchospasm or status asthmaticus? Yes No			
8. Will the beneficiary use Tezspire in combination with anti-IgE, anti-IL4, or anti-IL5 monoclonal antibody agents (e.g., benralizumab,			
omalizumab, mepolizumab, reslizumab, dupilumab)? Yes No			
9. Does the beneficiary have hypersensitivity to tezepelumab-ekko (Tezspire) or any of its excipients? Yes No			
10. Does the beneficiary have an active or untreated helminth infection? Yes No			
11. Will Tezspire be administered concurrently with live vaccines? Yes No			
For continuation of therapy, please answer questions 1-13 **Reauthorizations can be for up to 6 months **			
12. While on Tezspire, has the beneficiary experienced an improvement in asthma symptoms, asthma exacerbations, or airway function as			
evidenced by a decrease in ≥ 1 of the following? Yes No Please indicate which one(s):			
a. Use of systemic corticosteroids			
b. Two-fold or greater decrease in inhale	ed corticosteroid use for at least 3 days		
c. Hospitalizations			
d. ER visits			
e. Unscheduled visits to healthcare provider			
f. Improvement from baseline in FEV1			
13. Has the beneficiary experienced any serious treatment-related adverse events (e.g., parasitic [helminth] infection, severe hypersensitivity			
reactions)? YesNo ** Places provide modical records decumenting the honoficiary's current Asthma status and response to Tazenira treatment **			
** Please provide medical records documenting the beneficiary's current Asthma status and response to Tezspire treatment **			
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Signature of Prescriber:	Date	5:	

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of

material fact may subject me to civil or criminal liability.