



Tezspire

Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720.** If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect® 1-800-237-2767.

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Patient's Name: _____ **Date:** _____
Patient's ID: _____ **Patient's Date of Birth:** _____
Physician's Name: _____
Specialty: _____ **NPI#:** _____
Physician Office Telephone: _____ **Physician Office Fax:** _____

Referring Provider Info: ☐ Same as Requesting Provider

Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Rendering Provider Info: ☐ Same as Referring Provider ☐ Same as Requesting Provider

Name: _____ **NPI#:** _____
Fax: _____ **Phone:** _____

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Required Demographic Information:

Patient Weight: _____ *kg*

Patient Height: _____ *cm*

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

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Site of Service Questions:

- A. Where will this drug be administered?
- | | |
|---|---|
| <input type="checkbox"/> Ambulatory surgical, <i>skip to Clinical Questions</i> | <input type="checkbox"/> Home infusion, <i>skip to Clinical Questions</i> |
| <input type="checkbox"/> Off-campus Outpatient Hospital | <input type="checkbox"/> On-campus Outpatient Hospital |
| <input type="checkbox"/> Physician office, <i>skip to Clinical Questions</i> | <input type="checkbox"/> Pharmacy, <i>skip to Clinical Questions</i> |
- B. Is this request to continue previously established treatment with the requested medication?
- ☐ Yes - This is a continuation of an existing treatment.
- ☐ No - This is a new therapy request (patient has not received requested medication in the last 6 months). *skip to Clinical Criteria Questions*
- C. Has the patient experienced an adverse event with the requested product that has not responded to conventional interventions (eg acetaminophen, steroids, diphenhydramine, fluids, other pre-medications or slowing of the infusion rate) or a severe adverse event (anaphylaxis, anaphylactoid reactions, myocardial infarction, thromboembolism, or seizures) during or immediately after an infusion? ***ACTION REQUIRED: If Yes, Attach supporting clinical documentation.*** ☐ Yes, *skip to Clinical Criteria Questions* ☐ No
- D. 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? ***ACTION REQUIRED: If Yes, Attach supporting clinical documentation.*** ☐ Yes, *skip to Clinical Criteria Questions* ☐ No
- E. 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? ***ACTION REQUIRED: If Yes, Attach supporting clinical documentation.*** ☐ Yes ☐ No

Clinical Criteria Questions:

1. What is the diagnosis?
- ☐ Severe Asthma
- ☐ Other _____
2. What is the ICD-10 code? _____
3. Will the patient receive Tezspire as monotherapy (i.e., without any other asthma medications such as inhaled corticosteroids)? ☐ Yes ☐ No
4. Will the patient receive Tezspire concomitantly with other biologics indicated for asthma (e.g., Cinqair, Dupixent, Fasenna, Nucala, Xolair)? ☐ Yes ☐ No
5. Is this request for continuation of therapy with Tezspire? ☐ Yes ☐ No *If No, skip to #8*
6. Is the patient currently receiving Tezspire through samples or a manufacturer's patient assistance program? *If Yes or Unknown, skip to #8* ☐ Yes ☐ No ☐ Unknown
7. Has asthma control improved on Tezspire treatment as demonstrated by a reduction in the frequency and/or severity of symptoms and exacerbations OR a reduction in the daily maintenance oral corticosteroid dose? ***ACTION REQUIRED: If Yes, please attach supporting chart notes or medical record documentation of improved asthma control.*** ☐ Yes ☐ No
8. Does the patient have uncontrolled asthma as demonstrated by having one or more asthma exacerbation resulting in hospitalization or emergency medical care visit within the past year?
ACTION REQUIRED: If Yes, please attach supporting chart notes or medical records of asthma exacerbation(s) and skip to #11. ☐ Yes ☐ No
9. Does the patient have uncontrolled asthma as demonstrated by having two or more asthma exacerbations requiring oral or injectable corticosteroid treatment within the past year?
ACTION REQUIRED: If Yes, please attach supporting chart notes, medical records, or claims history of asthma exacerbations and previous corticosteroid treatment and skip to #11. ☐ Yes ☐ No

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- 10 Does the patient have uncontrolled asthma as demonstrated by having poor symptom control (frequent symptoms or reliever use, activity limited by asthma, night waking due to asthma) within the past year?

ACTION REQUIRED: If Yes, please attach supporting chart notes or medical records of symptom control.

☐ Yes ☐ No

11. Prior to receiving Tezspire, did the patient have inadequate asthma despite current treatment with both of the following medications at optimized doses? **ACTION REQUIRED: If Yes, please attach supporting chart notes, medical records, or claims history of previous medications tried including drug, dose, frequency, and duration.**

☐ Yes ☐ No

a) High dose inhaled corticosteroid

b) Additional controller (long-acting beta₂-agonist, long-acting muscarinic antagonist, leukotriene modifier, or sustained-release theophylline)

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

X _____

Prescriber or Authorized Signature

Date (mm/dd/yy)

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