

Service Authorization (SA) Form

Tezspire® (tezepelumab-ekko)

If the following information is not complete, correct, or legible, the SA process can be delayed.

Please use one form per member.

MEMBER INFORMATION

Last Name:

First Name:

Medicaid ID Number:

Date of Birth:

Weight in Kilograms:

PRESCRIBER INFORMATION

Last Name:

First Name:

NPI Number:

Phone Number:

Fax Number:

DRUG INFORMATION

Drug Name/Form:

Strength:

Dosing Frequency:

Length of Therapy:

Quantity per Day:

The Virginia Department of Medical Assistance Services considers the use of concomitant therapy with Cinqair®, Dupixent®, Fasenra®, Nucala®, Tezspire™ and Xolair® to be experimental and investigational. Safety and efficacy of these combinations have **NOT** been established and will **NOT** be permitted.

(Form continued on next page.)

Member's Last Name:

Member's First Name:

DIAGNOSIS AND MEDICAL INFORMATION

For severe* asthma initial approval, complete the following questions to receive a 6-month approval:

1. Is the member 12 years of age or older? **AND**

☐ Yes ☐ No

2. Does the member have a diagnosis of severe* asthma? **AND**

☐ Yes ☐ No

3. Will coadministration with another monoclonal antibody be avoided (e.g., omalizumab, mepolizumab, reslizumab, benralizumab, dupilumab)? **AND**

☐ Yes ☐ No

4. Will this be used for add-on maintenance treatment in members regularly receiving **both** (unless otherwise contraindicated) of the following:

- Medium- to high-dose inhaled corticosteroids; **AND**
- An additional controller medication (e.g., long-acting beta agonist, leukotriene modifiers)?

☐ Yes ☐ No

5. Has the member had two or more exacerbations in the previous year requiring oral or injectable corticosteroid treatment (in addition to the regular maintenance therapy defined above) **or** one exacerbation resulting in a hospitalization? **AND**

☐ Yes ☐ No

6. Does the member have at least one of the following for assessment of clinical status:

- Use of systemic corticosteroids
- Use of inhaled corticosteroids
- Number of hospitalizations, ER visits, or unscheduled visits to healthcare provider due to condition
- Forced expiratory volume in 1 second (FEV₁)? **AND**

☐ Yes ☐ No

(Form continued on next page.)

Member's Last Name:

Member's First Name:

7. Has the member tried and failed an adequate trial of the 2 different preferred products (Fasenra® and Xolair®)?

☐ Yes ☐ No ☐ N/A

If N/A was selected for question 7 please answer the following:

a. Does the member lack an eosinophilic phenotype with blood eosinophils ≥ 150 cells/ μ L? **AND**

☐ Yes ☐ No

b. Does the member lack a serum IgE level < 30 IU/mL? **OR**

☐ Yes ☐ No

c. Does the member have another predicted intolerance the preferred agents? (Answer below)

☐ Yes ☐ No

For severe* asthma renewal, complete the following questions to receive a 12-month approval:

8. Has the member been assessed for toxicity? **AND**

☐ Yes ☐ No

9. Does the member have improvement in asthma symptoms or asthma exacerbations as evidenced by decrease in one or more of the following:

- Use of systemic corticosteroids
- Hospitalizations
- ER visits
- Unscheduled visits to healthcare provider
- Improvement from baseline in forced expiratory volume in 1 second (FEV₁)?

☐ Yes ☐ No

***Components of severity for classifying asthma as severe may include any of the following (not all-inclusive):**

- Symptoms throughout the day
- Nighttime awakenings, often 7 times/week
- SABA use for symptom control occurs several times per day
- Extremely limited normal activities
- Lung function (percent predicted FEV₁) $< 60\%$
- Exacerbations requiring oral systemic corticosteroids are generally more frequent and intense relative to moderate asthma

(Form continued on next page.)

Member's Last Name:

Member's First Name:

Prescriber Signature (Required)

Date

By signature, the physician confirms the above information is accurate and verifiable by member records.

Please include ALL requested information; Incomplete forms will delay the SA process.

Submission of documentation does NOT guarantee coverage by the Department of Medical Assistance Services.

Fax this form to 1-866-940-7328

Pharmacy PA call center: 1-800-310-6826