

TEZSPIRE Federal Employee Program. PRIOR APPROVAL REQUEST

Additional information is required to process your claim for prescription drugs. Please complete the patient portion, and have the prescribing physician complete the

Service Benefit Plan **Prior Approval** P.O. Box 52080 MC 139 Phoenix, AZ 85072-2080

Send completed form to:

Attn. Clinical Services

Fax: 1-877-378-4727 **Provider Information** (required) Patient Information (required) Date: Provider Name: NPI: Patient Name: Specialty: Date of Birth: □Male ☐Female Office Phone: Office Fax: Sex: Office Street Address: Street Address: City: State: Zip: City: State: Zip: Patient ID: Physician Signature: PHYSICIAN COMPLETES

All approved requests are subject to review by a clinical specialist for final validation and coverage determination once all required documentation has been received. Current utilization, including samples, does not guarantee approval of coverage.

Tezspire

(tezepelumab-ekko)

**Check www.fepblue.org/formulary to confirm which medication is part of the patient's benefit

NOTE: Form must be completed in its entirety for processing
Is this request for brand or generic? □Brand □Generic
1. Will the patient need more than 3 units every 84 days? □Yes* □No
*If YES, please specify the requested quantity: units every 84 days
2. Has the patient been on this medication continuously for the last 6 months excluding samples? <i>Please select answer below:</i>
□ NO – this is INITIATION of therapy, please answer the following questions:
a. Does the patient have a diagnosis of severe asthma? □Yes □No
b. Has patient had inadequate control of asthma symptoms after a minimum of 3 months of compliant use defined as greater than or equal to 50 percent adherence with a corticosteroid inhaler in combination with a long acting beta2-agonist within the past 6 months? □Yes □No*
*If NO, has patient had inadequate control of asthma symptoms after a minimum of 3 months of compliant use defined as greater than or equal to 50 percent adherence with a corticosteroid inhaler in combination with a long acting muscarinic antagonist within the past 6 months? \(\textstyle{\textstyle{1}}\)Yes \(\textstyle{\textstyle{1}}\)No
c. Will this medication be used as add-on maintenance treatment? □Yes* □No
*If YES, please answer the following questions:
i. Will this medication be used in combination with a medium or high-dose inhaled corticosteroid? □Yes □No
ii. Will this medication be used in combination with an additional controller medication such as a long acting beta2 agonist or leukotriene modifier? Yes No
☐ YES – this is a PA renewal for CONTINUATION of therapy, please answer the following questions:
a. Does the patient have a diagnosis of asthma? □Yes □No
b. Has the patient had a documented decrease in exacerbations OR an improvement in symptoms? □Yes □No
c. Has the patient decreased utilization of rescue medications? □Yes □No
d. Has the patient been compliant on Tezspire therapy? □Yes □No
e. Will this medication be used as add-on maintenance treatment? □Yes □No
3. Will the patient be given live vaccines while on this therapy? □Yes □No
4. Will this medication be used for the relief of acute bronchospasm or status asthmaticus? □Yes □No
5. Will this medication be used in combination with another monoclonal antibody for the treatment of asthma or COPD? □Yes □No * <i>If YES</i> , please specify the medication:
PAGE 1 of 2 – Please fax back PAGES 1 and 2 with natient's medical records



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To ensure a quick and accurate response to your prior approval request, please **submit medical records** (**e.g.**, **chart notes**, **laboratory values**) pertaining to the diagnosis only. Please do not send in medical records of other diagnoses in order to streamline the process. Please use this page as a **guideline** of what documentation is required to process the prior authorization request.

*For more efficient processing, please provide the page number of the documented information in the medical record

Documentation Required:
□NOT used for the relief of acute bronchospasm or status asthmaticus PAGE of
□NO dual therapy with another monoclonal antibody PAGE of
□NOT given concurrently with live vaccines PAGE of
Documentation Required for <u>INITIATION</u> of Therapy:
□Severe asthma PAGE of
□ Inadequate control of symptoms after a minimum of 3 months of compliant use with greater than or equal to 50% adherence with ONE of the following within the past 6 months: PAGE of • Inhaled corticosteroids & long acting beta₂ agonist • Inhaled corticosteroids & long acting muscarinic antagonist □ Used as add-on maintenance treatment and will be receiving ALL of the following: PAGE of • Medium or high-dose inhaled corticosteroid • Additional controller medication (e.g., long acting beta₂ agonist, leukotriene modifier)
Documentation Required for <u>CONTINUATION</u> of Therapy:
□Decreased exacerbations OR improvement in symptoms PAGE of
□Decreased utilization of rescue medications PAGE of
□Compliant on therapy PAGE of
☐ Used as add-on maintenance treatment PAGE of

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