



Fasenra

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 (SOS):

- A. Indicate the site of service requested:
- | | |
|---|--|
| <input type="checkbox"/> On Campus Outpatient Hospital | <input type="checkbox"/> Off Campus Outpatient Hospital |
| <input type="checkbox"/> Home infusion, <i>skip to Criteria Questions</i> | <input type="checkbox"/> Physician office, <i>skip to Criteria Questions</i> |
| <input type="checkbox"/> Ambulatory surgical, <i>skip to Criteria Questions</i> | <input type="checkbox"/> Pharmacy, <i>skip to Criteria Questions.</i> |
- B. Which product is being requested?
- Fasenra
 Fasenra Pen *Skip to criteria questions*
- C. 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
- D. Has the patient experienced an adverse event with the requested product that has not responded to conventional interventions (eg acetaminophen, steroids, diphenhydramine, fluids, or other pre- medications) or a severe adverse event (anaphylaxis, anaphylactoid reactions, myocardial infarction, thromboembolism, or seizures) during or immediately after an infusion? **ACTION REQUIRED: If Yes, please attach supporting clinical documentation.**
- Yes, *skip to Clinical Criteria Questions* No
- E. 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, please attach supporting clinical documentation.
- Yes, *skip to Clinical Criteria Questions* No
- F. 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, please attach supporting clinical documentation. Yes No

Criteria Questions:

1. What is the diagnosis? Asthma Other, please specify _____
2. What is the ICD-10 code? _____
3. Is the medication prescribed by or in consultation with an allergist, immunologist, or pulmonologist?
 Yes No
4. Will the patient continue to use maintenance asthma treatments (e.g., inhaled corticosteroid, additional controller) in combination with Fasenra? Yes No
5. Will the patient receive Fasenra concomitantly with other biologics indicated for asthma (e.g., Cinqair, Dupixent, Nucala, Tezspire, Xolair)? Yes No
6. Is the request for continuation of therapy with Fasenra? Yes No *If No, skip to #10*
7. Is the patient currently receiving Fasenra through samples or a manufacturer’s patient assistance program?
If Yes or Unknown, skip to #10 Yes No Unknown
8. Has asthma control improved on Fasenra treatment as demonstrated by a reduction in the frequency and/or severity of symptoms and exacerbations? **ACTION REQUIRED: If Yes, please attach supporting chart notes or medical record documentation of improved asthma control.**
 Yes *If Yes, no further questions* No
9. Has asthma control improved on Fasenra treatment as demonstrated by a reduction in the daily maintenance oral corticosteroid dose? **ACTION REQUIRED:**

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If yes, please attach supporting chart notes or medical record documentation of improved asthma control.

Yes No *No further questions*

10. Does the patient have uncontrolled asthma as demonstrated by experiencing two or more asthma exacerbations requiring oral or injectable corticosteroid treatment within the past year? **ACTION REQUIRED: If yes, please submit supporting chart notes, medical records, or claims history of previous corticosteroid use for asthma exacerbations.** *If Yes, skip to #13* Yes No
11. Does the patient have uncontrolled asthma as demonstrated by experiencing one or more asthma exacerbation resulting in hospitalization or emergency medical care visit within the past year? **ACTION REQUIRED: If yes, please submit supporting chart notes, medical records of previous asthma exacerbations requiring hospitalization or emergency medical visit** *If Yes, skip to #13* Yes No
12. Does the patient have uncontrolled asthma as demonstrated by experiencing 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 submit supporting chart notes or medical records.** Yes No
13. Prior to receiving Fasenra, did the patient have inadequate asthma control despite current treatment with both of the following medications at optimized doses? a) High dose inhaled corticosteroid, b) additional controller (i.e., long acting beta2-agonist, long acting muscarinic antagonist, leukotriene modifier, or sustained-release theophylline). **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
14. What is the patient's baseline (e.g., before significant oral steroid use) blood eosinophil count in cells per microliter? **ACTION REQUIRED: Please attach supporting chart note(s) or medical record with the patient's baseline blood eosinophil count.** _____ cells per microliter Unknown
15. Is the patient dependent on systemic corticosteroids? **Action Required: Please attach supporting chart note(s) or medical record showing patient's dependance on systemic corticosteroids.** Yes No

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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