



Fasenra

CareFirst 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

What is the ICD-10 code? _____

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. Where will this drug be administered?
- ☐ Ambulatory surgical, *skip to Clinical Criteria Questions*
 - ☐ Home infusion, *skip to Clinical Criteria Questions*
 - ☐ Off-campus Outpatient Hospital, *Continue to B*
 - ☐ On-campus Outpatient Hospital, *Continue to B*
 - ☐ Physician office, *skip to Clinical Criteria Questions*
 - ☐ Pharmacy, *skip to Clinical Criteria Questions*
- B. Is the patient less than 14 years of age?
- ☐ Yes, *skip to Clinical Criteria Questions*
 - ☐ No, *Continue to C*
- C. Is this request to continue previously established treatment with the requested medication? **Action Required: If No, please attach supporting clinical documentation.**
- ☐ Yes - This is a continuation of an existing treatment., *Continue to D*
 - ☐ 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, *Continue to E*
- 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, *Continue to F*
- 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, *skip to Clinical Criteria Questions*
 - ☐ No, *Continue to G*
- G. Are **all** alternative infusion sites (pharmacy, physician office, ambulatory care, etc.) **greater than 30 miles** from the patient's home? **ACTION REQUIRED: If Yes, please attach supporting documentation.**
- ☐ Yes, *continue to Clinical Criteria Questions*
 - ☐ No, *continue to Clinical Criteria Questions*

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Criteria Questions:

1. Will the requested drug be used concomitantly with any other biologic or targeted synthetic drug for the same indication?

☐ Yes, *Continue to 2*

☐ No, *Continue to 2*

2. What is the diagnosis?

☐ Severe asthma, *Continue to 3*

☐ Other, please specify. _____, *Continue to 3*

3. Is the requested drug being prescribed by or in consultation with an allergist, immunologist, or pulmonologist?

☐ Yes, *Continue to 4*

☐ No, *Continue to 4*

4. Is the patient 6 years of age or older?

☐ Yes, *Continue to 5*

☐ No, *Continue to 5*

5. Is the request for continuation of therapy with the requested drug?

☐ Yes, *Continue to 6*

☐ No, *Continue to 11*

6. Is the patient currently receiving Fasenra through samples or a manufacturer's patient assistance program?

☐ Yes, *Continue to 11*

☐ No, *Continue to 7*

☐ Unknown, *Continue to 11*

7. Will the requested drug be used for the treatment of severe asthma?

☐ Yes, *Continue to 8*

☐ No, *Continue to 8*

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 chart notes or medical record documentation supporting improvement in asthma control.

☐ Yes, *Continue to 10*

☐ No, *Continue to 9*

9. Has asthma control improved on Fasenra treatment as demonstrated by a reduction in the daily maintenance oral corticosteroid dose? **ACTION REQUIRED:** If Yes, please attach chart notes or medical record documentation supporting improvement in asthma control.

☐ Yes, *Continue to 10*

☐ No, *Continue to 10*

10. Will the patient continue to use maintenance asthma treatments (e.g., inhaled corticosteroid, additional controller) in combination with Fasenra?

☐ Yes, *No Further Questions*

☐ No, *No Further Questions*

11. Has the patient received in the past year or is currently receiving a biologic drug (e.g., Dupixent, Nucala) indicated for treatment of severe asthma (excluding receiving the drug via samples or a manufacturer's patient assistance program)? **ACTION REQUIRED:** If Yes, please attach chart notes, medical record documentation, or claims history supporting previous medications tried including drug, dose, frequency and duration.

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- ☐ Yes, *Continue to 20*
☐ No, *Continue to 12*

12. Will the requested drug be used for the treatment of severe asthma?

- ☐ Yes, *Continue to 13*
☐ No, *Continue to 13*

13. 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 attach chart notes, medical record documentation, or claims history supporting previous corticosteroid use for asthma exacerbations including drug, dose, frequency and duration.

- ☐ Yes, *Continue to 16*
☐ No, *Continue to 14*

14. Does the patient have uncontrolled asthma as demonstrated by experiencing one or more asthma exacerbation(s) resulting in hospitalization or emergency medical care visit(s) within the past year? **ACTION REQUIRED:** If Yes, please attach chart notes or medical record documentation of previous asthma exacerbation(s) requiring hospitalization or emergency medical visit(s).

- ☐ Yes, *Continue to 16*
☐ No, *Continue to 15*

15. 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 attach chart notes or medical record documentation of poor symptom control.

- ☐ Yes, *Continue to 16*
☐ No, *Continue to 16*

16. Prior to requesting Fasenra, did the patient have inadequate asthma control despite current treatment with both of the following drugs at optimized doses: A) High-dose inhaled corticosteroid, AND 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 chart notes, medical record documentation, or claims history supporting previous medications tried including drug, dose, frequency and duration.

- ☐ Yes, *Continue to 17*
☐ No, *Continue to 17*

17. Prior to requesting Fasenra, what is the patient's baseline (e.g., before significant oral steroid use) blood eosinophil count in cells per microliter? Indicate baseline blood eosinophil count in cells per microliter. **ACTION REQUIRED:** Please attach chart notes or medical record documentation with the patient's baseline blood eosinophil count.

- ☐ Greater than or equal to 150 cells per microliter _____ cells/microliter **ACTION**

REQUIRED: *Submit supporting documentation, Continue to 19*

- ☐ Less than 150 cells per microliter _____ cells/microliter **ACTION REQUIRED:** *Submit supporting documentation, Continue to 18*

- ☐ Unknown, *Continue to 18*

18. Is the patient dependent on systemic corticosteroids? **ACTION REQUIRED:** If Yes, please attach chart notes or medical record documentation showing patient's dependence on systemic corticosteroids.

- ☐ Yes, *Continue to 19*
☐ No, *Continue to 19*

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19. Will the patient continue to use maintenance asthma treatments (e.g., inhaled corticosteroid, additional controller) in combination with Fasenra?

☐ Yes, *Continue to 20*

☐ No, *Continue to 20*

20. Is a loading dose prescribed?

☐ Yes, *Continue to 21*

☐ No, *No Further Questions*

21. Does the prescribed loading dose exceed a dose of 30 mg?

☐ Yes, *Continue to 22*

☐ No, *Continue to 22*

22. What is the prescribed loading dose?

☐ 10 mg, *No further questions*

☐ 30 mg, *No further questions*

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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