



PRIOR AUTHORIZATION REQUEST

FASENRA

Patient Information:

Name:	
Member ID:	
Address:	
City, State, Zip:	
Date of Birth:	

Prescriber Information:

Name:	
NPI:	
Phone Number:	
Fax Number:	
Address:	
City, State, Zip:	

Requested Medication

Rx Name:	
Rx Strength:	
Rx Quantity:	
Rx Frequency:	
Rx Route of Administration:	
Diagnosis and ICD Code:	

Your patient's prescription benefit requires that we review certain requests for coverage with the prescriber. You have prescribed a medication for your patient that requires Prior Authorization before benefit coverage or coverage of additional quantities can be provided. Please complete the following questions then fax this form to the toll-free number listed below. Upon receipt of the completed form, prescription benefit coverage will be determined based on the plan's rules.

SECTION A: Please note that supporting clinical documentation is required for ALL PA requests. Pharmacy prior authorization reviews can be subject to trial with additional medications that are not listed within the criteria. The policies are subject to change based on COMAR requirements, MDH transmittals and updates to treatment guidelines.

1 Is the request an INITIAL or CONTINUATION of therapy?

Initial (If checked, go to 8)

Continuation (If checked, go to 2)

2 Will the requested medication be used in combination with other monoclonal antibodies? Yes No

[Note: Examples of monoclonal antibody therapies include benralizumab,

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mepolizumab and dupilumab.]
 [If yes, no further questions.]

3	Is the patient currently receiving the requested medication? [If no, skip to question 8.]	Yes	No
4	Has the patient been established on therapy for at least 3 months? [If no, skip to question 8.]	Yes	No
5	Has the patient been receiving medication samples for the requested medication? [If yes, skip to question 8.]	Yes	No
6	Does the patient have a previously approved prior authorization (PA) on file with the current plan? [Note: If the patient does NOT have a previously approved PA on file for the requested medication with the current plan, the renewal request will be considered under initial therapy.] [If no, skip to question 8.]	Yes	No
7	Has documentation been submitted to confirm that the patient has experienced a clinically significant response, as determined by the provider? ACTION REQUIRED: Submit supporting documentation. [No further questions.]	Yes	No
8	Will the requested medication be used in combination with other monoclonal antibodies? [Note: Examples of monoclonal antibody therapies include benralizumab, mepolizumab and dupilumab.] [If yes, no further questions.]	Yes	No
9	What is the diagnosis or indication? <input type="checkbox"/> Asthma (If checked, go to 10) <input type="checkbox"/> Eosinophilic Granulomatosis with Polyangiitis (EGPA) [formerly known as Churg-Strauss Syndrome] (If checked, go to 26) <input type="checkbox"/> Chronic Obstructive Pulmonary Disease (COPD) (If checked, no further questions) <input type="checkbox"/> Hypereosinophilic Syndrome (HES) (If checked, no further questions) <input type="checkbox"/> Other (If checked, no further questions)		
10	Is the patient 6 years of age or older? [If no, no further questions.]	Yes	No
11	Does the patient have a blood eosinophil count of GREATER THAN OR EQUAL TO 150 cells per microliter within the previous 6 weeks or within 6 weeks prior to treatment with any anti-interleukin (IL)-5 therapy?	Yes	No

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[Note: Examples of anti-IL-5 therapies include Fasenra, Nucala, and Cinqair.]
 [If no, no further questions.]

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|----|--|-----|----|
| 12 | Has the patient received at least 3 consecutive months of combination therapy with an inhaled corticosteroid (ICS)?
[Note: Examples of ICS include Aerospa n , Alvesco, ArmonAir RespiClick, Arnuity Ellipta, Asmanex Twisthaler/hydrofluroalkane (HFA), Flovent Diskus/HFA, Pulmicort Flexhaler, Qvar/Qvar RediHaler, and budesonide suspension for inhalation (Pulmicort Respules, generics).]
[If no, skip to question 15.] | Yes | No |
| 13 | Has the patient received at least 3 consecutive months of combination therapy with at least ONE additional asthma controller/maintenance medication?
[Note: Examples of additional asthma controller/maintenance medications include long-acting beta2-agonist (LABA) (such as Serevent Diskus), inhaled long-acting muscarinic antagonist (LAMA) (such as Spiriva Respimat), leukotriene receptor antagonists (such as montelukast tablets/granules [Singulair, generics], zafirlukast tablets [Accolate, generics]), and theophylline (such as Theo 24, TheoChron ER, generics).]
[If yes, skip to question 16.] | Yes | No |
| 14 | Has the patient already received anti-interleukin (IL)-5 therapy (such as Cinqair, Fasenra, or Nucala) used concomitantly with an inhaled corticosteroid (ICS) for at least 3 consecutive months instead of a trial with one additional asthma controller/maintenance medication?
[Note: Examples of ICS include Aerospa n , Alvesco, ArmonAir RespiClick, Arnuity Ellipta, Asmanex Twisthaler/hydrofluroalkane (HFA), Flovent Diskus/HFA, Pulmicort Flexhaler, Qvar/Qvar RediHaler, and budesonide suspension for inhalation (Pulmicort Respules, generics).]
[If yes, skip to question 16.] | Yes | No |
| 15 | Has the patient received at least 3 consecutive months of a combination inhaler containing BOTH an inhaled corticosteroid (ICS) and a long-acting beta2-agonist (LABA) instead of receiving therapy with both an ICS and one additional asthma controller/maintenance medication?
[Note: Examples of combination ICS/LABA inhalers include Advair Diskus (generic Wixela Inhub; authorized generics), Advair hydrofluroalkane (HFA), AirDuo RespiClick (authorized generics), Breo Ellipta, Dulera, and Symbicort].
[If no, no further questions.] | Yes | No |
| 16 | Will the requested medication be used in combination with an inhaled corticosteroid (ICS) or ICS-containing combination inhaler?
[If no, no further questions.] | Yes | No |
| 17 | Is the patient's asthma uncontrolled as defined by the patient experiencing TWO OR MORE asthma exacerbations requiring treatment with systemic corticosteroids in the previous year?
[If yes, skip to question 22.] | Yes | No |

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18	Is the patient's asthma uncontrolled as defined by the patient experiencing ONE asthma exacerbation requiring hospitalization in the previous year? [If yes, skip to question 22.]	Yes	No
19	Is the patient's asthma uncontrolled as defined by a forced expiratory volume in 1 second (FEV1) LESS THAN 80% predicted? [If yes, skip to question 22.]	Yes	No
20	Is the patient's asthma uncontrolled as defined by a forced expiratory volume in 1 second (FEV1)/forced vital capacity (FVC) LESS THAN 0.80? [If yes, skip to question 22.]	Yes	No
21	Does the patient's asthma worsen upon tapering of oral corticosteroid therapy? [If no, no further questions.]	Yes	No
22	Has documentation been submitted to confirm that the patient has had an intolerance, contraindication to, or failed treatment for at least 3 months with preferred interleukin (IL)-4 and interleukin (IL)-13 receptor inhibitor, Dupixent? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
23	Has documentation been submitted to confirm that the patient has had an intolerance, contraindication to, or failed treatment for at least 3 months with interleukin (IL)-5 inhibitor, Nucala? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]	Yes	No
24	Is the requested medication prescribed by or in consultation with an allergist, immunologist, or pulmonologist? [If no, no further questions.]	Yes	No
25	Does the dose of the requested medication exceed the Food and Drug Administration (FDA) approved label dosing for the indication? [Note: Dosing: 12 years old and older: SUBQ: 30 mg every 4 weeks for the first 3 doses, then once every 8 weeks. Dosing Children: 6 to less than 12 years old AND if weight less than 35 kg: SUBQ: 10 mg every 4 weeks for the first 3 doses, then once every 8 weeks OR if weight greater than or equal to 35 kg: SUBQ: 30 mg every 4 weeks for the first 3 doses, then once every 8 weeks.] [No further questions.]	Yes	No
26	Is the patient 18 years of age or older? [If no, no further questions.]	Yes	No
27	Has the patient tried a minimum of 4 weeks of therapy with a corticosteroid (for example, prednisone)? [If no, no further questions.]	Yes	No

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28	<p>Does the patient have/or had a blood eosinophil level of 150 cells per microliter or GREATER within the previous 6 weeks or within 6 weeks prior to treatment with any anti-interleukin (IL)-5 therapy? [Note: Examples of anti-IL-5 therapies include Nucala, Cinqair, and Fasenra.] [If no, no further questions.]</p>	Yes	No
29	<p>Does the patient have active, non-severe disease? [Note: Non-severe disease is defined as vasculitis without life- or organ-threatening manifestations and examples of symptoms in patients with non-severe disease include rhinosinusitis, asthma, mild systemic symptoms, uncomplicated cutaneous disease, and mild inflammatory arthritis.] [If no, no further questions.]</p>	Yes	No
30	<p>Has documentation been submitted to confirm that the patient has had an intolerance, contraindication to, or failed treatment for at least 3 months with interleukin (IL)-5 inhibitor, Nucala? ACTION REQUIRED: Submit supporting documentation. [If no, no further questions.]</p>	Yes	No
31	<p>Is the requested medication being prescribed by or in consultation with an allergist, immunologist, pulmonologist or a rheumatologist? [If no, no further questions.]</p>	Yes	No
32	<p>Does the dose of the requested medication exceed the Food and Drug Administration (FDA) approved label dosing for the indication? [Note: Dosing: 30 mg every 4 weeks.]</p>	Yes	No

Please document the diagnoses, symptoms, and/or any other information important to this review:

SECTION B: Physician Signature

PHYSICIAN SIGNATURE	DATE
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FAX COMPLETED FORM TO: 1-833-896-0656

Disclaimer: An authorization is not a guarantee of payment. Member must be eligible at the time services are rendered. Services must be a covered Health Plan Benefit and medically necessary with prior authorization as per Plan policy and procedures.

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