

## State of Oklahoma SoonerCare

## Fasenra® (Benralizumab) Prior Authorization Form

Member Name:	Date of Birth:	Member ID#:
Drug Information		
□Physician billing (HCPCS code:) □Pharmacy billing (NDC:)		
Dose: Regimen:	· · · · · · · · · · · · · · · · · · ·	Start Date:
Billing Provider Information		
Provider NPI: Provider Name:		
Provider Phone: Provider Fax:		
Prescriber Information		
Prescriber NPI: Prescriber Name:		
Prescriber Phone: Prescriber Phone:	escriber Fax:	Specialty:
<b>Criteria</b>		
information must be provided and SoonerCare may verify through further requested documentation. The member's drug history will be reviewed prior to approval. <i>Initial approvals will be for the duration of six months.</i> 1. What is the diagnosis for which the medication is being prescribed?  2. Severe eosinophilic phenotype asthma  3. Other, please list:  2. Will benralizumab be used as add-on maintenance treatment for severe eosinophilic phenotype asthma?		
Yes No 3. If yes, please indicate member's daily med Drug/Dose:	Drug/Dos	se:
4. Baseline blood eosinophil count:	Date Dete	ermined:
<ol> <li>Has the member been evaluated by an allergist, pulmonologist, or pulmonary specialist within the last 12 months (or an advanced care practitioner with a supervising physician who is an allergist, pulmonologist, or pulmonary specialist)? Yes No</li> <li>If yes, please include name of specialist:</li> <li>Is member compliant with a medium-to-high-dose inhaled corticosteroid (ICS) plus at least 1 additional controller medication? Yes No</li> <li>Does member require daily systemic corticosteroids despite compliant use of a medium-to-high-dose ICS plus at least 1 additional controller medication? Yes No</li> <li>If answer is 'no' to previous question, please list number and dates of exacerbations requiring systemic corticosteroids within lost 12 months: Number: Dates of exacerbations:</li> </ol>		
corticosteroids within last 12 months: Number: Dates of exacerbations: 10. Please check all that apply:		
<ul> <li>Member has failed a medium-to-high-dose ICS used compliantly within the last 3-6 consecutive months.</li> <li>Drug/Dose:</li> <li>Member has failed at least 1 other asthma controller medication used in addition to the medium-to-high-dose ICS compliantly for at least the past 3 months.</li> <li>Drug/Dose:</li> </ul>		
<ul> <li>11. For Fasenra® prefilled syringe, will it be administered in a health care setting by a health care professional prepared to manage anaphylaxis? Yes No</li> <li>12. For Fasenra® prefilled autoinjector pen, has member or caregiver been trained by a health care professional on subcutaneous administration, monitoring for any allergic reactions, and storage of Fasenra® prefilled autoinjector pen? Yes No</li> </ul>		
Members must be adherent for continued approval. Compliance will be evaluated for continued approval.		
Prescriber Signature:  (By signature, the physician confirms the criteria information above is accurate and verifiable in patient records.)  Please do not send in chart notes. Specific information/documentation will be requested if necessary. Failure to		
complete this form in full will result in processing delays.		

## PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy Pharmacy Management Consultants Product Based Prior Authorization Unit

> Fax: 1-800-224-4014 Phone: 1-800-522-0114 Option 4

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