



Would you like to use electronic prior authorization? Consider using Surescripts, our electronic prior authorization portal at providerportal.surescripts.net/ProviderPortal/login OR fax completed prior authorization request form to 800-750-9692.

All requested data must be provided. **Incomplete forms or forms without the chart notes will be returned**

Pharmacy Coverage Guidelines are available at sentarahealthplans.com/en/providers/authorizations/prescription-drugs

Dupixent (Medicare)

REQUIRED: Office notes, labs, and medical testing relevant to request showing medical justification to support diagnosis					
Member Information					
Member Name (first & last):		Date of Birth:	Gender: <input type="checkbox"/> Male <input type="checkbox"/> Female	Height:	
Member ID:	City:		State: Weight:		
Prescribing Provider Information					
Requestor's Name:		Requestor's Phone Number:		Requestor's Fax Number:	
Provider Name (first & last):	Specialty	NPI:		DEA:	
Office Address:		City:		State: Zip Code:	
Office Contact:		Office Phone:		Office Fax:	
Dispensing Provider/Pharmacy Information					
Place of Administration:	<input type="checkbox"/> Self-Administered <input type="checkbox"/> Physician's Office <input type="checkbox"/> Home <input type="checkbox"/> Home Infusion Center <input type="checkbox"/> Outpatient Infusion Center Name: _____				
Agency NPI:	Agency Name:		Agency Phone Number:		
Agency Address			Agency Fax Number:		
City:		State:		Zip:	
Dispensing Location:	<input type="checkbox"/> Retail Pharmacy <input type="checkbox"/> Mail Order <input type="checkbox"/> Physician's Office <input type="checkbox"/> Specialty Pharmacy <input type="checkbox"/> Other				
Pharmacy Name:		Pharmacy Phone:		Pharmacy Fax:	
Pharmacy NPI:					
Requested Medication Information					
Medication request is NOT for an FDA approved, or compendia-supported diagnosis (check one): <input type="checkbox"/> Yes <input type="checkbox"/> No			Diagnosis:	ICD-10 Code:	
Are there any contraindications to formulary medications? If yes, please specify:			Is this a New Request or Continuation of Therapy: <input type="checkbox"/> New, start date: ____/____/_____ <input type="checkbox"/> Continuation, date of last treatment: ____/____/_____		
Directions for Use:			Strength:	Dosage Form	
			Duration:	Quantity:	Days Supply:
What medication(s) has the member tried and failed for this diagnosis? Please specify below including duration of treatment.					
Turn-Around Time for Review:					
<input type="checkbox"/> Standard <input type="checkbox"/> Urgent: Waiting standard time for decision could seriously harm life, health, or ability to regain maximum function, you can ask for an expedited decision.					
Signature: _____					



Sentara®

Health Plans Member First Name: _____ Member Last Name: _____

Member ID: _____ Member Date of Birth: _____

Clinical Information:

* Indicate questions that are required to be answered

Q1. Please select applicable diagnosis: *

- Chronic Obstructive Pulmonary Disease (COPD)
- Chronic Spontaneous Urticaria (CSU)
- Eosinophilic Esophagitis (EOE)
- Prurigo Nodularis (PN)
- Atopic Dermatitis (AD)
- Asthma
- Chronic Rhinosinusitis with Nasal Polyps (CRSwNP)
- Other

Q2. For COPD: Which of the following has the member met?

- Labs submitted showing a blood eosinophils at least 300 cells per microliter taken within previous 6 weeks or prior to monoclonal antibody therapy
- FEV1/FVC ratio less than 0.7 and FEV1 at least 30% but not more than 80% post-bronchodilator
- Currently treated with triple or dual therapy (long-acting beta-2 agonist (LABA)/long-acting muscarinic antagonists (LAMA)/ inhaled corticosteroid (ICS) or ong-acting muscarinic antagonists (LAMA)/long-acting beta-2 agonist (LABA)) for at least 3 months
- Has signs or symptoms of chronic bronchitis for at least 3 months in the previous 12 months

Q3. For COPD: Has the member experienced COPD exacerbations in the past 12 months meeting ONE of the following?

- 2 or more episodes requiring treatment with oral corticosteroids or antibiotics
- Episode required hospitalization

Q4. For COPD: Will the member continue using dual or triple therapy while on requested medication?

- Yes
- No

Q5. For CSU: Does the member have documentation of the following?

- Urticaria present for no less than 6 weeks
- Member experiences spontaneous occurrence of wheals or hives that are pruritic in nature, not painful, more than 3 days per week
- Symptoms persist despite taking doses of second generation H1-antihistamines (e.g. cetirizine) AND a leukotriene receptor antagonist (i.e. montelukast) for at least two weeks

Q6. For EOE: Has diagnosis been confirmed by endoscopic biopsy showing at least 15 intraepithelial eosinophils per hpf?

- Yes
- No

Q7. For EOE: Have secondary causes have been ruled out?



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Yes

No

Q8. For EOE: Has the member tried and failed 8 weeks therapy with a prescription strength Proton Pump Inhibitor (i.e. omeprazole 40 mg)?

Yes

No

Q9. For PN: Does the member have at least 20 nodular lesions and has experienced 6 weeks of pruritus?

Yes

No

Q10. For Reauthorization: Has the member responded positively to therapy as determined by the prescribing physician?

Yes

No

Q11. For Asthma: Which of the following has the member met?

Member has a blood eosinophil at least 150 cells per microliter w/in prior 6 weeks OR has oral corticosteroid-dependent asthma

Member has received both an inhaled corticosteroid AND an adjunct maintenance medication (i.e. montelukast)
(Note: Exception to the requirement for a trial of an asthma maintenance medication can be made if already established on monoclonal antibodies used concomitantly w/an inhaled corticosteroid)

Asthma is uncontrolled defined by member experiencing one of the following (A, B, C, D, or E): A) 2 or more asthma exacerbations requiring treatment with systemic corticosteroid in last 12 months, B) 1 or more asthma exacerbations requiring hospital or emergency department visit in last 12 months, C) FEV1 less than 80% predicted, D) FEV1/FVC less than 0.80, OR E) asthma worsens with tapering of oral corticosteroid.

Q12. For Asthma: Will the member continue to receive treatment with 1 inhaled corticosteroid or 1 inhaled corticosteroid-containing combo inhaler?

Yes

No

Q13. For CRSwNP: Is the member receiving treatment with an intranasal corticosteroid and experiencing rhinosinusitis symptoms like nasal obstruction, rhinorrhea, or reduction/loss of smell?

Yes

No

Q14. For CRSwNP: Has the member received treatment within previous 2 yrs or has contraindication to systemic corticosteroid treatment OR if member had prior surgery for nasal polyps?

Yes

No

Q15. For CRSwNP: Will the member continue to receive treatment with an intranasal corticosteroid while on the requested medication?



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Yes

No