

Specialty Medication Prior Authorization Cover Sheet

(This cover sheet should be submitted along with a Pharmacy Prior Authorization Medication Fax Request Form. Please refer to www.uhcprovider.com for medication fax request forms.)

Patient Information

Patient's Name:

Insurance ID:

Date of Birth:

Height:

Weight:

Address:

Apartment #:

City:

State:

Zip Code:

Phone Number:

Alternate Phone:

Sex: Male

Female

Provider Information

Provider's Name:

Provider ID Number:

Address:

City:

State:

Zip Code:

Suite Number:

Building Number:

Phone Number:

Fax number:

Provider's Specialty:

Medication Information

Medication:

Quantity:

ICD10 Code:

Directions:

Diagnosis:

Refills:

Physician Signature:**

Initial here if DAW:

Physician Signature:** *By signing above, the physician is providing the specialty pharmacy with a prescription that can be used to facilitate the dispensing and/or coordination of delivery for the requested medication.*

Medication Instructions

Has the patient been instructed on how to **Self-Administer**?

Yes No

Is this medication a **New Start**?

Yes No

If continuation please provide the following: Initiation Date: / / Date of Last Dose: / /

Is there documentation of positive clinical response to current therapy? Yes No

****Please attach any pertinent clinical information that would pertain to support stated diagnosis.**

Additional clinical information may be needed depending on your patients plan, including medication(s) previously tried and failed.

Delivery Instructions

Note: Delivery coordination requires a **"Physician Signature"** above and complete
"Provider Information" and **"Patient Information"**

Note: All necessary ancillary supplies are provided free of charge to the patient at the time of delivery

Ship to: Physician's Office Patient's Address Date medication is needed: / /

Medication Administered: Home Health Self-Administered LTC Physician's Office

PRIOR AUTHORIZATION REQUEST FORM

Please complete this entire form and fax it to: **866-940-7328**. If you have questions, please call **800-310-6826**.

This form contains multiple pages. Please complete all pages to avoid a delay in our decision.
Allow at least 24 hours for review.

Section A – Member Information				
First Name:	Last Name:		Member ID:	
Address:				
City:	State:		ZIP Code:	
Phone:	DOB:		Allergies:	
Primary Insurance Information:				
Is the requested medication <input type="checkbox"/> New or <input type="checkbox"/> Continuation of Therapy? If continuation, list start date: _____				
Is this patient currently hospitalized? <input type="checkbox"/> Yes <input type="checkbox"/> No If recently discharged, list discharge date: _____				
Section B - Provider Information				
First Name:	Last Name:			M.D./D.O.
Address:	City:		State:	ZIP code:
Phone:	Fax:	NPI #:		Specialty:
Office Contact Name / Fax attention to:				
Section C - Medical Information				
Medication:				Strength:
Directions for use:				Quantity:
Diagnosis (Please be specific & provide as much information as possible):				ICD-10 CODE:
Is this member pregnant? <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, what is this member's due date? _____				
Section D – Previous Medication Trials				
Medications	Strength	Directions	Dates of Therapy	Reason for failure / discontinuation
Section E – Additional information and Explanation of why preferred medications would not meet the patient's needs: Please refer to the patient's PDL for a list of preferred alternatives				

Member First name:	Member Last name:	Member DOB:
Clinical and Drug Specific Information		
ALL REQUESTS		
<input type="checkbox"/> Yes <input type="checkbox"/> No	Does the patient have one of the following diagnoses? <i>(If yes, check which applies)</i> <ul style="list-style-type: none"> <input type="checkbox"/> Moderate chronic atopic dermatitis <input type="checkbox"/> Severe chronic atopic dermatitis <input type="checkbox"/> Moderate to severe Asthma 	
<input type="checkbox"/> Yes <input type="checkbox"/> No	Is the patient receiving Dupixent in combination with any of the following? <i>(If yes, check which applies)</i> <ul style="list-style-type: none"> <input type="checkbox"/> Anti-interleukin-5 therapy [e.g. Nucala (mepolizumab), Cinqair (reslizumab), Fasenra (benralizumab)] <input type="checkbox"/> Anti-IgE therapy [e.g. Xolair (omalizumab)] <input type="checkbox"/> Biologic medication [e.g. Enbrel (etanercept), Rituxan (rituximab), Remicade/Inflectra (infliximab)] <p><i>If yes, list rationale:</i></p>	
<input type="checkbox"/> Yes <input type="checkbox"/> No	Is Dupixent prescribed by one of the following? <i>(If yes, check which applies)</i> <ul style="list-style-type: none"> <input type="checkbox"/> Dermatologist <input type="checkbox"/> Allergist <input type="checkbox"/> Immunologist <input type="checkbox"/> Pulmonologist 	
CHRONIC ATOPIC DERMATITIS		
<input type="checkbox"/> Yes <input type="checkbox"/> No	Did the patient require systemic immunosuppressive therapy for control with any of the following? <i>(If yes, check which applies and complete Section D above)</i> <ul style="list-style-type: none"> <input type="checkbox"/> Cyclosporine <input type="checkbox"/> Azathioprine <input type="checkbox"/> Methotrexate <input type="checkbox"/> Mycophenolate mofetil 	
<input type="checkbox"/> Yes <input type="checkbox"/> No	Does the patient have a history of failure, contraindication, or intolerance to any of the following topical therapies? <i>(If yes, check which applies and complete Section D above)</i> <ul style="list-style-type: none"> <input type="checkbox"/> Medium to very-high potency topical corticosteroid [e.g., Elocon (mometasone furoate), Synalar (fluocinolone acetonide), Lidex (fluocinonide)] <input type="checkbox"/> Topical calcineurin inhibitor [e.g., Elidel (pimecrolimus), Protopic (tacrolimus)]. <input type="checkbox"/> Eucrisa (crisaborole) 	
MODERATE TO SEVERE ASTHMA (continued on next page)		
<input type="checkbox"/> Yes <input type="checkbox"/> No	Is the patient's asthma uncontrolled or inadequately controlled as defined by at least one of the following? <i>(If yes, check which applies)</i> <ul style="list-style-type: none"> <input type="checkbox"/> Poor symptom control (e.g., Asthma Control Questionnaire [ACQ] score consistently greater than 1.5 or Asthma Control Test [ACT] score consistently less than 20) <p><i>List rationale:</i></p> <ul style="list-style-type: none"> <input type="checkbox"/> Two or more bursts of systemic corticosteroids for at least 3 days each in the previous 12 months <p><i>List corticosteroid and dates:</i></p> <ul style="list-style-type: none"> <input type="checkbox"/> Asthma-related emergency treatment (e.g., emergency room visit, hospital admission, or unscheduled physician's office visit for nebulizer or other urgent treatment) <p><i>List emergency treatment:</i></p> <ul style="list-style-type: none"> <input type="checkbox"/> Airflow limitation (e.g., after appropriate bronchodilator withhold forced expiratory volume in 1 second [FEV1] less than 80% predicted [in the face of reduced FEV1-forced vital capacity [FVC] defined as less than the lower limit of normal]) <p><i>List rationale:</i></p> <ul style="list-style-type: none"> <input type="checkbox"/> Patient is currently dependent on oral corticosteroids for the treatment of asthma <p><i>List oral corticosteroid:</i></p> 	
<input type="checkbox"/> Yes <input type="checkbox"/> No	Is there submission of medical records (e.g., chart notes, laboratory values, etc.) documenting that asthma is an eosinophilic phenotype as defined by a baseline (pre-dupilumab treatment) peripheral blood eosinophil level ≥ 150 cells/μL within the past 6 weeks? <p><i>If yes, list peripheral blood eosinophil level and date:</i></p>	

Member First name:		Member Last name:	Member DOB:
<input type="checkbox"/> Yes <input type="checkbox"/> No	Is the patient currently dependent on oral corticosteroids for the treatment of asthma? <i>If yes, list corticosteroids:</i>		
<input type="checkbox"/> Yes <input type="checkbox"/> No	Will Dupixent be used in combination with one high dose (appropriately adjusted for age) combination inhaled corticosteroid (ICS)/long-acting beta2 agonist (LABA)? <i>If yes, list combo ICS/LABA product:</i>		
<input type="checkbox"/> Yes <input type="checkbox"/> No	Will Dupixent be used in combination with one high-dose (appropriately adjusted for age) ICS product? <i>If yes, list high dose ICS product:</i>		
<input type="checkbox"/> Yes <input type="checkbox"/> No	Will Dupixent be used in combination with any additional asthma controller medications? (e.g., long-acting beta2 agonist, theophylline, leukotriene receptor antagonist) <i>If yes, list additional product:</i>		
CONTINUATION OF THERAPY			
<input type="checkbox"/> Yes <input type="checkbox"/> No	Does the patient have a documented positive clinical response to Dupixent as demonstrated by any of the following? (If yes, check which applies) <input type="checkbox"/> Reduction in the frequency of exacerbations <input type="checkbox"/> Decreased utilization of rescue medications <input type="checkbox"/> Increase in percent predicted FEV1 from pretreatment baseline <input type="checkbox"/> Reduction in severity or frequency of asthma-related symptoms (e.g., wheezing, shortness of breath, coughing, etc.) <input type="checkbox"/> Reduction in oral corticosteroid requirements <i>If other, list response:</i>		

Physician Signature: _____ **Date:** _____

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