

PRESCRIPTION DRUG MEDICATION REQUEST FORM FAX TO 1-866-240-8123

DUPIXENT PRIOR AUTHORIZATION FORM PATIENT INFORMATION						
Subscribe	r ID Number	PATIENTIN	FORMATION	Group Number		
Patient Name			Patient Telephone Number	Date of Birth		
Patient Address			City	tate Zip Code		
PRESCRIBER INFORMATION						
Physician Name			Phone	Fax		
Physician Address			City	State Zip Code		
Suite / Building Physician Signature		Physician Signature		Date		
		MEDICATION	INFORMATION			
Reque	sted Strength:					
☐ 100mg/0.67ml Syringe ☐ 200mg/1.14ml Syringe			☐ 300mg/2ml S	yringe		
		☐ 200mg/1.14ml Pen	☐ 300mg/2ml P	en		
Quantity: Number of pens/syringes Does the patient require induction dosing of 4 pens/syringes for the first 4 weeks of therapy? □ Yes □ No						
Diagno	osis:					
		CLINICAL	. CRITERIA			
If Dupi	xent is being used to trea	t moderate-to-severe atopic	dermatitis, please answ	er the following:		
1. Dupixent is being prescribed by a:						
	☐ Dermatologist	☐ Allergist	☐ Immunologist	☐ Other:		
2.	Does the patient have a ☐ Yes ☐ No	topic dermatitis with facial or	anogenital involvement?			
3.	3. Has the patient experienced therapeutic failure, intolerance, or contraindication to any of the following? Please select ALL that apply:					
	 ☐ A topical corticosteroid (e.g. Betamethasone, Clobetasol, Triamcinolone, etc.) ☐ Topical Tacrolimus (Protopic) ☐ Topical Pimecrolimus (Elidel) 					
4.	 Does the patient have severe atopic dermatitis with a large BSA (body surface area) which would make topical therapy impractical to apply? ☐ Yes ☐ No 					
5.	. Does the patient have severe atopic dermatitis with severely damaged skin? ☐ Yes ☐ No					
6.	Is this a request for read ☐ Yes ☐ No	uthorization?				
	a. If YES , has the pation ☐ Yes ☐ No	ent experienced positive clini	ical response to therapy v	vith Dupixent?		

If Dupix	kent is being used to treat moderate-to-severe <u>asthma</u> , please answer the following:			
1.	Please provide ALL of the following:			
	a. Patient's pretreatment FEV1:% predicted			
2.	Does the patient have FEV1 reversibility of at least 12% and 200 milliliters (ml) after albuterol (salbutamol) administration? ☐ Yes ☐ No			
3.	Does the patient have eosinophilic phenotype with blood eosinophil count greater than or equal to 150 cells/microliter? □ Yes □ No			
4.	Is the patient currently taking daily or alternate-day oral corticosteroids? $\hfill \Box$ Yes $\hfill \Box$ No			
5.	Is the patient using a medium- or high-dose inhaled corticosteroid? ☐ Yes ☐ No			
6.	Is the patient using a long-acting beta agonist? ☐ Yes ☐ No			
7.	Is this a request for reauthorization? ☐ Yes ☐ No			
	 a. If YES, please select ALL that apply:			
If Dupix	cent is being used to treat chronic rhinosinusitis with nasal polyposis , please answer the following:			
1.	Please provide:			
	 a. Patient's baseline bilateral nasal polyp score (from 0 to 8): The Nasal Polyp Score, the sum of right and left nostril scores, is used to characterize the patient's polyps. Each nostril is scored on a scale of 0 to 4, with the total score being the sum of left and right nostril scores. 0 = no polyps 4 = severe disease with large polyps causing complete obstruction of the inferior nasal cavity 			
	 b. Patient's baseline nasal congestion score (from 0 to 3): The Nasal Congestion Score is a tool used to measure changes in nasal congestion and obstruction. 0 = no symptoms 3 = severe symptoms 			
2.	Has the patient experienced therapeutic failure, intolerance, or contraindication to the following: Please select ALL that apply: An intranasal corticosteroid A 14-day course of oral corticosteroids			
3.	Is this a request for reauthorization? Yes No a. If YES, please select ALL that apply: Patient has a decrease in the nasal polyp score Patient has a reduction in the nasal congestion/obstruction severity score			

If Dupixent is being used to treat eosinophilic esophagitis , please answer the following:					
1.	Does the patient weigh at least 40 kg? ☐ Yes ☐ No				
2.	Does the patient have an esophageal eosinophil count greater than or equal to 15 eos/hpf (eosinophils/high power field) on esophageal biopsy? ☐ Yes ☐ No				
3.	Does the patient have clinical symptoms of esophageal dysfunction (e.g., dysphagia, food impaction, gastroesophageal reflux)? \Box Yes \Box No				
4.	Has the patient experienced two or more episodes of dysphagia per week? ☐ Yes ☐ No				
5.	Has the patient experienced therapeutic failure, contraindication, or intolerance to high-dose proton-pump inhibitor (PPI) therapy (e.g. omeprazole or pantoprazole 80 mg/day)? ☐ Yes ☐ No				
6.	Is this a request for reauthorization? Yes No If YES, please select ALL that apply: Patient experienced histological remission (i.e. less than 15 eos/hpf) on esophageal biopsy Patient experienced reduced severity or frequency of dysphagia Patient experienced reduced severity or frequency of clinical symptoms of esophageal dysfunction (e.g. food impaction, gastroesophageal reflux)				

The submitting provider certifies that the information provided is true, accurate, and complete and the requested services are medically indicated and necessary to the health of the patient. Note: Payment is subject to member eligibility. Authorization does not guarantee payment.

INSTRUCTIONS FOR COMPLETING THIS FORM

- 1. Submit a separate form for each medication.
- 2. Complete **ALL** information on the form.

NOTE: The prescribing physician (PCP or Specialist) should, in most cases, complete the form.

- 3. Please provide the physician address as it is required for physician notification.
- 4. Fax the **completed** form and all clinical documentation to **1-866-240-8123**

Or mail the form to: Clinical Services,

120 Fifth Avenue, MC PAPHM-043B, Pittsburgh, PA 15222