

**Xolair® Prior Authorization Request Form (Page 1 of 2)**

<b>Member Information (required)</b>			<b>Provider Information (required)</b>		
Member Name:			Provider Name:		
Insurance ID#:			NPI#:		Specialty:
Date of Birth:			Office Phone:		
Street Address:			Office Fax:		
City:	State:	Zip:	Office Street Address:		
Phone:			City:	State:	Zip:
<b>Medication Information (required)</b>					
Medication Name:			Strength:		Dosage Form:
<input type="checkbox"/> Check if requesting <b>brand</b>			Directions for Use:		
<input type="checkbox"/> Check if request is for <b>continuation of therapy</b>					
<b>Clinical Information (required)</b>					
<b>Select the diagnosis below:</b> <input type="checkbox"/> Allergic asthma <input type="checkbox"/> Chronic idiopathic urticaria <input type="checkbox"/> Other diagnosis: _____ ICD-10 Code(s): _____					
<b>Provider's Specialty:</b> Select if Xolair is prescribed by or in consultation with one of the following specialists: <input type="checkbox"/> Allergist/immunologist <input type="checkbox"/> Dermatologist <input type="checkbox"/> Pulmonologist					
<b>For allergic asthma, answer the following:</b> Does the patient have a diagnosis of moderate to severe persistent allergic asthma? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have a pre-treatment serum immune globulin (IgE) level between 30 to 1300 IU/mL? <input type="checkbox"/> Yes <input type="checkbox"/> No Does the patient have positive skin test or in vitro reactivity to a perennial aeroallergen? <input type="checkbox"/> Yes <input type="checkbox"/> No Are symptoms adequately controlled on a high-dose inhaled corticosteroid and a long-acting beta2-agonist combination for at least 3 months? <input type="checkbox"/> Yes <input type="checkbox"/> No Has the patient been adherent within a 12 month period and is currently adherent with asthma therapy? <input type="checkbox"/> Yes <input type="checkbox"/> No					
<b>Reauthorization:</b> Select if the patient has experienced improvement with treatment as defined by one of the following: <input type="checkbox"/> Reduction in number of asthma exacerbations from baseline (i.e., asthma exacerbation requiring treatment with systemic corticosteroids or doubling of inhaled corticosteroid [ISC] dose from baseline) <input type="checkbox"/> Improvement in forced expiratory volume in 1 second (FEV1) from baseline <input type="checkbox"/> Decreased use of rescue medications from baseline					
<b>For chronic idiopathic urticaria, answer the following:</b> Does the patient have persistent symptoms (itching and hives) for at least 4 consecutive weeks despite titrating to an optimal dose with a second generation H1 antihistamine? <input type="checkbox"/> Yes <input type="checkbox"/> No Select if the patient has had inadequate response or intolerance to the following additional therapies: <input type="checkbox"/> Doxepin <input type="checkbox"/> Hydroxyzine <input type="checkbox"/> H1 antihistamine <input type="checkbox"/> Leukotriene receptor antagonist <input type="checkbox"/> H2 antagonist					
<b>&lt;continued on next page&gt;</b>					

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**Reauthorization:**

Has the patient's disease status been re-evaluated since the last authorization to confirm the patient's condition warrants continued treatment?  Yes  No

Has the patient experienced a reduction in itching severity or a reduction in the number of hives from baseline?  Yes  No

**Are there any other comments, diagnoses, symptoms, medications tried or failed, and/or any other information the physician feels is important to this review?**

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Please note:

This request may be denied unless all required information is received.

**Please fax this form to 1-844-712-8129 to initiate a prior authorization review for the member and medication above.**

**Please note: plan benefits may limit or exclude coverage of specific medications including those requested on this form.**

I certify, to the best of my knowledge, the statements and information provided on this form are factual and correct.

Provider/Representative (and Title): \_\_\_\_\_ Date: \_\_\_\_\_

**PROACT INTERNAL USE ONLY:**

**Clinical Review Decision**

Approved, through

Denied (documentation attached, if necessary)

**Tracking:**

1 <sup>st</sup> Attempt		2 <sup>nd</sup> Attempt		Letter Mailed:	
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