

XOLAIR (omalizumab) (preferred) PRIOR AUTHORIZATION FORM

Prior authorization guidelines for **Monoclonal Antibodies, Anti-IL, Anti-IgE and Quantity Limits/Daily Dose Limits** are available on the DHS Pharmacy Services website at <http://www.dhs.pa.gov/provider/pharmacyservices/index.htm>.

PRIOR AUTHORIZATION INFORMATION		PRESCRIBER INFORMATION	
<input type="checkbox"/> New request	<input type="checkbox"/> Renewal request	Total # of pages: _____	Prescriber name:
Name of office contact:		Specialty:	
Contact's phone number:		State license #:	
LTC facility contact/phone:		NPI:	MA Provider ID#:
BENEFICIARY INFORMATION		Street address:	
Beneficiary name:		Suite #:	City/state/zip:
Beneficiary ID#:	DOB:	Phone:	Fax:

CLINICAL INFORMATION

Medication requested:	<input type="checkbox"/> Xolair 150 mg/ml syringe	<input type="checkbox"/> Xolair 75 mg/0.5 ml syringe
	<input type="checkbox"/> Xolair 150 mg vial	<input type="checkbox"/> Xolair _____
Dose/directions:	Quantity:	Duration: _____ months
Diagnosis:	Dx code (<i>required</i>):	Weight: _____ lbs / kg
Specialty Pharmacy Drug Program: Which specialty pharmacy will be used? <input type="checkbox"/> Diplomat Specialty <input type="checkbox"/> Walgreens Specialty		

Initial Requests

1. Is Xolair being prescribed by or in consultation with a specialist?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Submit documentation of consultation, if applicable.
2. For a diagnosis of asthma: Is the beneficiary being treated for moderate to severe persistent asthma induced by an unavoidable perennial allergen (pollen, mold, dust mites, etc) and inadequately controlled by inhaled corticosteroids?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Submit documentation, including results of allergen reactivity test.
3. For a diagnosis of asthma: Does the beneficiary have a serum total IgE measurement between 30 international units (IU)/ml and 1300 IU/ml?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Submit documentation of test results.
4. For a diagnosis of asthma: Is the beneficiary currently receiving optimally titrated doses, or have a contraindication or intolerance to, any of the following? <input type="checkbox"/> inhaled glucocorticoid <input type="checkbox"/> long-acting beta-agonist (LABA) <input type="checkbox"/> leukotriene modifier <input type="checkbox"/> other (eg, tiotropium, theophylline): _____	<input type="checkbox"/> Yes <input type="checkbox"/> No	Submit documentation of medication regimen and response.
5. For a diagnosis of chronic idiopathic urticaria (CIU): Does the beneficiary have a history of urticaria for a period of ≥ 3 months?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Submit documentation.
6. For a diagnosis of CIU: Does the beneficiary require the use of steroids to control urticarial symptoms?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Submit documentation.
7. For a diagnosis of CIU: Does the beneficiary have a history of trial and failure, contraindication, or intolerance of all of the following at maximal tolerated doses? <i>Check all that apply.</i> <input type="checkbox"/> H ₁ antihistamine <input type="checkbox"/> leukotriene modifier <input type="checkbox"/> H ₂ antihistamine <input type="checkbox"/> dapsone, sulfasalazine, or hydroxychloroquine	<input type="checkbox"/> Yes <input type="checkbox"/> No	Submit documentation.
8. Will the beneficiary be monitored and/or treated for helminth infection as recommended in package labeling?	<input type="checkbox"/> Yes <input type="checkbox"/> No	

Renewal Requests

1. For a diagnosis of asthma, has the beneficiary experienced measurable evidence of improvement in asthma severity?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Submit documentation of response to therapy.
2. For a diagnosis of chronic idiopathic urticaria, does the beneficiary have documentation of improvement in symptoms and rationale for continued use of Xolair?	<input type="checkbox"/> Yes <input type="checkbox"/> No	Submit documentation of response to therapy and rationale for continued use.
3. Will the beneficiary be monitored and/or treated for helminth infection as recommended in package labeling?	<input type="checkbox"/> Yes <input type="checkbox"/> No	

PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO DHS – PHARMACY DIVISION

Prescriber Signature:	Date:
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