



PRIOR AUTHORIZATION REQUEST FORM
CHRONIC SPONTANEOUS URTICARIA

Dupixent®, Xolair®

For authorization, please answer each question and fax this form PLUS chart notes back to Real Rx at 385-425-4052.

Failure to submit clinical documentation to support this request will result in a dismissal of the request.

If you have prior authorization questions, please call for assistance 385-425-5094.

Disclaimer: Prior authorization request forms are subject to change in accordance with Federal and State notice requirements.

Date:	Member Name:	ID#:
DOB:	Gender:	Physician:
Office Phone:	Office Fax:	Office Contact:
Height/Weight:	HCPCS Code:	

Member must try formulary preferred drugs before a request for a non-preferred drug may be considered. If treatment with preferred products has not been successful, you must submit which preferred products have been tried, dates of treatment, and reason for failure. Reasons for failure must meet the Health Plan medical necessity criteria.

Preferred/Non-Preferred

1. Preferred
 - a. Xolair® (omalizumab)
2. Non-Preferred
 - a. Dupixent® (dupilumab)

Dosing/Frequency: _____

Note: for the treatment of nasal polyps see Chronic Rhinosinusitis with Nasal Polyposis (CRSwNP)

If the request is for reauthorization, proceed to reauthorization section

Questions	Yes	No	Comments/Notes
1. Is the requested medication being purchased by the provider's office and to be billed under the medical benefit ('buy-and-bill')?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Is this request for an expedited review? By checking the "Yes" box to request an expedited review (24 hours), you are certifying that applying the standard review time frame (72 hours) may place the member's life, health, or ability to regain maximum function in serious jeopardy.	<input type="checkbox"/>	<input type="checkbox"/>	
3. Has the provider performed a medical evaluation that rules out other possible causes of urticaria?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
4. Has the member had a trial and failure of an H1-antihistamine at up to four times standard dosing used in combination with an H2-antihistamine?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
5. Has the member had a trial and failure of an H1-antihistamine used in combination with a leukotriene receptor antagonist or cyclosporine?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation

6. Is the request for dose escalation of Xolair®?	<input type="checkbox"/>	<input type="checkbox"/>	
7. For Dupixent®, does the member have a contraindication or intolerance to Xolair®?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
REAUTHORIZATION			
1. Is the request for reauthorization of therapy?	<input type="checkbox"/>	<input type="checkbox"/>	
2. Does clinical documentation show continued medical necessity and that the treatment has stabilized or improved the member's condition?	<input type="checkbox"/>	<input type="checkbox"/>	Please provide documentation
What medications and/or treatment modalities have been tried in the past for this condition? Please document name of treatment, reason for failure, treatment dates, etc.			
Additional information:			
Physician's Signature:			

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Policy: PHARM-164
 Origination Date: 06/11/2025
 Reviewed/Revised Date: 06/11/2025
 Next Review Date: 06/11/2026
 Current Effective Date: 07/01/2025

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