

Follow these steps to submit prior authorization requests when prescribing drugs covered under the medical benefit for Blue Cross Blue Shield of Michigan and Blue Care Network commercial members.

Michigan prescribers

To submit prior authorization requests electronically, first register for Availity® Essentials, our provider portal; refer to the [Register for web tools](#) page at bcbsm.com for details. Then:

1. Log in to [availity.com](#)*.
2. Click *Payer Spaces* on the menu bar and click the BCBSM and BCN logo.
3. On the Applications tab, click the tile for the appropriate NovoLogix web tool.
4. Within NovoLogix, click the *Authorizations* menu and select *Create Authorization*.
5. Enter the member's details and select the correct member on the contract.
6. Complete the required fields. This includes selecting the correct drug in the "Authorization Lines" section.
7. Click *Submit*, complete the protocol questions and click *Done*.

If you're registered for Availity but are not able to access it, submit your prior authorization request using the *Medication Authorization Request Form*, or MARF, that's on the next page.

Non-Michigan prescribers

When submitting a prior authorization request for the first time, prescribers located outside of Michigan should complete and submit:

- The *Medication Authorization Request Form*, or MARF, that's on the next page
- The [Application for access to NovoLogix for non-Michigan prescribers](#)

Submit these documents to the fax number or address that's on the MARF. Once we approve the request for access, we'll provide information about how to access the NovoLogix tool so that you can submit subsequent prior authorization requests electronically.

Note: Access to NovoLogix is available only to registered users. You must include a valid Type 1 (individual) NPI on the application for access to NovoLogix.

Information about NovoLogix

For more information about the NovoLogix web tool, look under the Training Resources heading on these webpages:

- [Blue Cross Medical-Benefit Drugs](#)
- [BCN Medical-Benefit Drugs](#)

If you need help with the NovoLogix tool, contact the Web Support Help Desk at 1-877-258-3932.

*Clicking this link means that you're leaving the Blue Cross Blue Shield of Michigan and Blue Care Network website. While we recommend this site, we're not responsible for its content.

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Blue Cross Blue Shield/Blue Care Network of Michigan
Medication Authorization Request Form
Xolair® (omalizumab) HCPCS CODE: J2357



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This form is to be used by participating physicians to obtain coverage for Xolair®. For commercial members only, please complete this form and submit via fax to 1-877-325-5979. If you have any questions regarding this process, please contact BCBSM Provider Relations and Servicing or the Medical Drug Helpdesk at 1-800-437-3803 for assistance.

PATIENT INFORMATION	PHYSICIAN INFORMATION
Name	Name
ID Number	Specialty
D.O.B. <input type="checkbox"/> Male <input type="checkbox"/> Female	Address
Pt weight (in kg) Date recorded: _____	City /State/Zip
Diagnosis	Phone/Fax: P: () - F: () -
Drug Name	NPI
Dose and Quantity	Contact Person
Directions	Contact Person/ Phone Ext.
Date of Service(s)	

STEP 1: DISEASE STATE INFORMATION

- Is this for Initiation or Continuation of therapy? ☐ Initiation ☐ Continuation Date patient started therapy: _____
- How is this medication being administered? ☐ Self-administered (**Please fax this completed form to BCBSM at (866) 601-4425**)
☐ Healthcare professional administered (**Continue to #3**)
- Will the patient receive the first 3 doses under the guidance of a health care provider? ☐ Yes ☐ No Comment: _____
- Site of administration? ☐ Provider office/Home infusion ☐ Other: _____
☐ Hospital outpatient facility (go to #4) Reason for Hospital Outpatient administration: _____
- Please specify location of administration if hospital outpatient infusion:** _____
- Please provide the NPI number for the place of administration:** _____
- Initiation AND Continuation of therapy:**
 - Will the patient be using Xolair in combination with other biologic agents (for example: Nucala, Fasenra, Cinqair or Dupixent) or targeted DMARD medications?
☐ Yes ☐ No Comment: _____
 - Please check the patient's diagnosis:** ☐ Moderate to Severe Allergic Asthma (AA, go to c then d) ☐ Nasal polyps (go to c then i)
☐ Chronic idiopathic urticaria (CIU, go to f) ☐ IgE-mediated food allergy (go to c, then k) ☐ Other _____
 - What is the patient's IgE level at the start of therapy? _____ IU/mL Date: _____
 - AA:** Which of the following tests did the patient receive for the diagnosis of moderate to severe allergic asthma?
☐ Positive skin test to a perennial aeroallergen (allergens with year-round exposure which may include molds, dust mites, cock roaches, animal feathers or dander, etc.)
☐ In-vitro reactivity to a perennial aeroallergen (allergens with year-round exposure which may include molds, dust mites, cock roaches, animal feathers or dander, etc.)
☐ N/A ☐ Other: _____
 - AA:** Which treatment(s) did not adequately control the patient's severe allergic asthma symptoms after a trial of at least 3 months?
☐ Systemic corticosteroid: _____ Date: Start: _____ End: _____
☐ High dose inhaled corticosteroids: _____ Date: Start: _____ End: _____
☐ Long acting beta2-agonist: _____ Date: Start: _____ End: _____
☐ Leukotriene receptor antagonist: _____ Date: Start: _____ End: _____
☐ Combination asthma inhaler with a HIGH dose corticosteroid and a long acting beta agonist: _____ Date: Start: _____ End: _____
☐ Combination asthma inhaler with a MEDIUM dose corticosteroid and a long acting beta agonist: _____ Date: Start: _____ End: _____
☐ Long acting muscarinic antagonist (LAMA): _____ Date: Start: _____ End: _____
☐ Other: _____ Date: Start: _____ End: _____
 - Chronic Idiopathic Urticaria (CIU):** How long has the patient been experiencing hives and itching (occurring daily or almost daily) in weeks?
☐ ≥ 6 weeks ☐ < 6 weeks ☐ Other: _____
 - CIU:** Have other diagnoses (such as Atopic Dermatitis, Contact Dermatitis, and reversible triggers) been ruled out?
☐ Yes ☐ No Comment: _____

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- h. **CIU:** Which medications did the patient try and experience treatment failure at maximally tolerated doses for at least 2 months?
- ☐ 1st Generation Antihistamine drug and dose (such as Benadryl): _____ **Start:** _____ **End:** _____
- ☐ 2nd Generation Antihistamine drug and dose (such as: Zyrtec, Claritin, Allegra): _____ **Start:** _____ **End:** _____
- ☐ H2 antagonist drug and dose (such as: Zantac or Pepcid): _____ **Start:** _____ **End:** _____
- ☐ Leukotriene receptor antagonist (such as: Singulair): _____ **Start:** _____ **End:** _____
- ☐ Hydroxyzine **Start:** _____ **End:** _____
- ☐ Doxepin **Start:** _____ **End:** _____
- ☐ Other: _____
- i. **Nasal polyps:** Is the patient currently receiving and will continue to receive a standard of care regimen for their diagnosis with Xolair?
- ☐ Yes ☐ No **Comment:** _____
- j. **Nasal polyps:** Has the patient tried and failed intranasal corticosteroids?
- ☐ Yes ☐ No **Comment:** _____
- k. **IgE-mediated food allergy:** Do the patient have a history of an allergic reaction following the consumption of peanuts, milk, eggs, wheat, cashews, hazelnuts or walnuts?
- ☐ Yes, please specify: _____ ☐ No **Comment:** _____
- l. **IgE-mediated food allergy:** Does the patient have a food allergy been confirmed by either:
- ☐ IgE specific antibodies, please specify IgE level (kUA/L): _____
- ☐ Food-specific skin prick test (SPT)
- m. **IgE-mediated food allergy:** Will the patient be on allergen avoidant diet while on Xolair?
- ☐ Yes ☐ No **Comment:** _____
- n. **IgE-mediated food allergy:** Does the patient have an active prescription and access to an epinephrine auto-injector?
- ☐ Yes ☐ No **Comment:** _____
- o. **IgE-mediated food allergy:** Will the patient be on any other food allergy desensitization treatments?
- ☐ Yes ☐ No **Comment:** _____

8. **Continuation request** (please answer above questions as well): **Xolair start date:** _____

- a. Have the patient's signs and symptoms improved with Xolair?
- ☐ Yes ☐ No, **Comment:** _____ ☐ Other: _____
- b. Please provide reason(s) why the patient needs to continue to have Xolair administered by a healthcare professional:
- ☐ Prior history of anaphylaxis including to Xolair, or other agents such as foods, drugs, or biologics
- ☐ Hypersensitivity reactions during the first 3 doses under the guidance of a healthcare provider
- ☐ Patient or caregivers who have been trained and are unable to recognize or treat symptoms of anaphylaxis
- ☐ Patient has co-morbidities or chronic medical conditions (such as: rheumatoid arthritis, Parkinson's disease), please specify: _____
- ☐ Other: _____

Please add any other supporting medical information necessary for our review

Coverage will not be provided if the prescribing physician's signature and date are not reflected on this document.

☐ Request for expedited review: I certify that applying the standard review time frame may seriously jeopardize the life or health of the member or the member's ability to regain maximum function

Physician's Name	Physician Signature	Date
Step 2: Checklist <input type="checkbox"/> Form Completely Filled Out <input type="checkbox"/> Attached Chart Notes		<input type="checkbox"/> Attach Diagnostic Tests
Step 3: Submit	By Fax: BCBSM Specialty Pharmacy Mailbox 1-877-325-5979	By Mail: BCBSM Specialty Pharmacy Program P.O. Box 312320, Detroit, MI 48231-2320