



Instructions:

This form is used by Kaiser Permanente and/or participating providers for coverage of **Xolair (omalizumab)**. Please complete and fax this form back to Kaiser Permanente within 24 hours [fax: [1-866-331-2104](tel:1-866-331-2104)]. If you have any questions or concerns, please call [1-866-331-2103](tel:1-866-331-2103). **Requests will not be considered unless all sections are complete.**

KP-MAS Formulary can be found at: [Pharmacy | Community Provider Portal | Kaiser Permanente](#)

1 – Patient Information

Patient Name: _____ Kaiser Medical ID#: _____ Date of Birth: _____

2 – Prescriber Information

Prescriber Name: _____ Specialty: _____ NPI: _____

Prescriber Address: _____

Prescriber Phone #: _____ Prescriber Fax #: _____

3 – Pharmacy Information

Pharmacy Name: _____ Pharmacy NPI: _____

Pharmacy Phone # _____ Pharmacy Fax #: _____

4 – Drug Therapy Requested

Drug 1: Name/Strength/Formulation: _____

Sig: _____

Drug 2: Name/Strength/Formulation: _____

Sig: _____

5– Diagnosis/Clinical Criteria

1. Is this request for initial or continuing therapy?

Initial therapy Continuing therapy, state start date: _____

2. Indicate the patient’s diagnosis for the requested medication: _____

Clinical Criteria:**If using for asthma:**

1. Prescriber is a Pulmonologist or Allergist,
 No Yes

2. AND member has diagnosis of uncontrolled moderate to severe asthma defined as any of the following:
 - a. ≥ 2 exacerbations in the past 12 months requiring systemic corticosteroids for more than 3 days
 - b. ≥ 1 asthma exacerbation(s) leading to hospitalization in the past 12 months
 - c. Dependence on daily oral corticosteroids (OCS) for asthma control
 - d. Poor symptom control (ACT score less than 20) No Yes

3. AND member has uncontrolled asthma despite good adherence (at least 75% over the past 3 months) to a regimen containing: a high dose inhaled corticosteroid, long-acting beta 2 agonist, AND long-acting muscarinic antagonist, and consideration given to use of a leukotriene receptor antagonist,
 No Yes

4. AND member is ≥ 6 years,
 No Yes

5. AND clinical diagnosis of allergic asthma,
 No Yes

6. AND if requiring Xolair q2week dosing, patient has documented treatment failure, contraindication, or inadequate response to Dupixent,
 No Yes

7. AND Xolair will NOT be used with Fasentra (benralizumab), Cinqair (reslizumab), Nucala (mepolizumab), Dupixent (dupilumab), or Tezspire (tezepelumab-ekko)
 No Yes

If using for nasal polyps:

1. Prescriber is an Allergist or ENT Specialist,
 No Yes

2. AND member has diagnosis of rhinosinusitis with nasal polyps,
 No Yes

3. AND member has history of failure, inadequate response, contraindication, or intolerance to Dupixent (dupilumab)
 No Yes

If using for chronic spontaneous urticaria:

1. Prescriber is an Allergist or Dermatologist,
 No Yes

2. AND member has diagnosis of chronic spontaneous urticaria,
 No Yes

3. AND member is 12 years of age or older,
 No Yes

4. AND member has tried and failed therapy for a minimum of 4 weeks on ALL of the following, unless contraindicated:
- a. At least two different high-dose second generation H1-antihistamines (e.g. loratadine, cetirizine) 2-4 times normal dose daily OR two second-generation H1-antihistamines in combination (e.g. fexofenadine 180 mg daily in the morning plus cetirizine 10-20 mg daily at bedtime),
 - b. AND montelukast in combination with a high-dose second generation H1-antihistamine,
 - c. AND H2-antihistamines (e.g. famotidine, ranitidine) in combination with a high-dose second generation H1-antihistamine

No Yes

For continuation of therapy, please respond to additional questions below:

- 1. Member has documentation of positive clinical response to Xolair therapy since last review
 No Yes
- 2. AND member continues to be under the care of a specialist
 No Yes

7 – Prescriber Sign-Off

Additional Information –

- 1. **Please submit chart notes/medical records for the patient that are applicable to this request.**
- 2. **If member has not tried preferred agent(s) please provide rationale/explanation and any additional supporting information that should be taken into consideration for the requested medication:**

I certify that the information provided is accurate. Supporting documentation is available for State audits.

Prescriber Signature:	Date:
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