

SPECIALTY GUIDELINE MANAGEMENT

XOLAIR (omalizumab)

POLICY

I. INDICATIONS

The indications below including FDA-approved indications and compendial uses are considered a covered benefit provided that all the approval criteria are met and the member has no exclusions to the prescribed therapy.

A. FDA-Approved Indications

1. Allergic asthma

Xolair is indicated for patients 6 years of age and older with moderate to severe persistent asthma who have a positive skin test or in vitro reactivity to a perennial aeroallergen and whose symptoms are inadequately controlled with inhaled corticosteroids.

Limitations of use: Xolair is not indicated for the relief of acute bronchospasm or status asthmaticus, or for treatment of other allergic conditions.

2. Chronic rhinosinusitis with nasal polyps (CRSwNP)

Xolair is indicated for add-on maintenance treatment of chronic rhinosinusitis with nasal polyps (CRSwNP) in adult patients 18 years of age and older with inadequate response to nasal corticosteroids.

3. IgE-mediated food allergy

Xolair is indicated for the reduction of allergic reactions (Type 1), including anaphylaxis, that may occur with accidental exposure to one or more foods in adult and pediatric patients aged 1 year and older with IgE-mediated food allergy.

Xolair is to be used in conjunction with food allergen avoidance.

Limitations of use: Xolair is not indicated for the emergency treatment of allergic reactions, including anaphylaxis.

4. Chronic spontaneous urticaria (CSU)

Xolair is indicated for the treatment of adults and adolescents 12 years of age and older with chronic spontaneous urticaria (CSU) who remain symptomatic despite H1 antihistamine treatment.

Limitations of use: Xolair is not indicated for treatment of other forms of urticaria.

B. Compendial Uses

1. Immune checkpoint inhibitor-related toxicities
2. Systemic mastocytosis

All other indications are considered experimental/investigational and not medically necessary.

II. DOCUMENTATION

Reference number(s)
1656-A

Submission of the following information is necessary to initiate the prior authorization review:

- A. Asthma:
 - 1. Initial requests:
 - i. Chart notes or medical record documentation showing pre-treatment IgE level.
 - ii. Chart notes, medical record documentation, or claims history supporting previous medications tried.
 - 2. Continuation requests: Chart notes or medical record documentation supporting improvement in asthma control.
- B. CRSwNP:
 - 1. Initial requests:
 - i. Chart notes or medical record documentation showing nasal endoscopy, anterior rhinoscopy, or computed tomography (CT) details (e.g., polyps location, size), or Meltzer Clinical Score or endoscopic nasal polyp score (NPS) (where applicable).
 - ii. Chart notes, medical record documentation, or claims history supporting previous medications tried. If therapy is not advisable, documentation of clinical reason to avoid therapy.
 - 2. Continuation requests: Chart notes or medical record documentation supporting positive response to therapy.
- C. IgE-mediated food allergy:
 - 1. Initial requests: Chart notes, medical record documentation, or laboratory tests showing the following (if applicable):
 - i. Pre-treatment allergen-specific IgE level
 - ii. Skin-prick test wheal diameter
 - iii. Pre-treatment serum IgE level
 - iv. Positive result of a physician controlled oral food challenge
 - v. History of a systemic reaction to a food
 - 2. Continuation requests: Chart notes or medical record documentation supporting positive response to therapy (e.g., decrease in hypersensitivity to food-allergen).
- D. CSU:
 - 1. Initial requests: Chart notes, medical record documentation, or claims history supporting previous medications tried showing an inadequate treatment response to a second-generation H1 antihistamine.
 - 2. Continuation requests: Chart notes or medical record documentation supporting response to therapy.
- E. Immune checkpoint inhibitor-related toxicity (initial requests): Chart notes or medical record documentation showing pre-treatment IgE level.
- F. Systemic mastocytosis (initial requests):
 - 1. Chart notes or medical record documentation supporting diagnosis of systemic mastocytosis.
 - 2. Chart notes, medical record documentation, or claims history of prerequisite therapies (if applicable).

III. PRESCRIBER SPECIALTIES

This medication must be prescribed by or in consultation with one of the following:

- A. Asthma: allergist/immunologist or pulmonologist
- B. CRSwNP: allergist/immunologist or otolaryngologist
- C. IgE-mediated food allergy: allergist/immunologist
- D. Chronic spontaneous urticaria: allergist/immunologist or dermatologist

IV. CRITERIA FOR INITIAL APPROVAL

A. Asthma

- 1. Authorization of 6 months may be granted for members 6 years of age or older who have previously received a biologic drug (e.g., Nucala, Cinqair) indicated for asthma.

2. Authorization of 6 months may be granted for treatment of moderate-to-severe asthma when all of the following criteria are met:
 - i. Member is 6 years of age or older.
 - ii. Member has a positive skin test or in vitro reactivity to at least one perennial aeroallergen.
 - iii. Member has a pre-treatment IgE level greater than or equal to 30 IU/mL.
 - iv. Member has uncontrolled asthma as demonstrated by experiencing at least one of the following within the past year:
 - a. Two or more asthma exacerbations requiring oral or injectable corticosteroid treatment.
 - b. One or more asthma exacerbation resulting in hospitalization or emergency medical care visit.
 - c. Poor symptom control (frequent symptoms or reliever use, activity limited by asthma, night waking due to asthma).
 - v. Member has inadequate asthma control despite current treatment with both of the following medications at optimized doses:
 - a. Medium-to-high-dose inhaled corticosteroid
 - b. Additional controller (i.e., long acting beta₂-agonist, long-acting muscarinic antagonist, leukotriene modifier, or sustained-release theophylline)
 - vi. Member will continue to use maintenance asthma treatments (e.g., inhaled corticosteroid, additional controller) in combination with the requested medication.

B. Chronic rhinosinusitis with nasal polyps (CRSwNP)

1. Authorization of 6 months may be granted for adult members who have previously received a biologic drug (e.g., Nucala, Dupixent) indicated for chronic rhinosinusitis with nasal polyps (CRSwNP).
2. Authorization of 6 months may be granted for treatment of CRSwNP when all of the following criteria are met:
 - i. Member is 18 years of age or older.
 - ii. Member has bilateral nasal polyps and chronic symptoms of sinusitis despite intranasal corticosteroid treatment for at least 2 months unless contraindicated or not tolerated.
 - iii. Member has one of the following:
 - a. A bilateral nasal endoscopy, anterior rhinoscopy, or computed tomography (CT) showing polyps reaching below the lower border of the middle turbinate or beyond in each nostril
 - b. Meltzer Clinical Score of 2 or higher in both nostrils
 - c. A total endoscopic nasal polyp score (NPS) of at least 5 with a minimum score of 2 for each nostril
 - iv. Member has symptoms of nasal blockage, congestion or obstruction plus one of the following additional symptoms:
 - a. Rhinorrhea (anterior/posterior)
 - b. Reduction or loss of smell
 - c. Facial pain or pressure
 - v. Member will continue to use a daily intranasal corticosteroid while being treated with the requested medication, unless contraindicated or not tolerated.

C. IgE-mediated food allergy

Authorization of 6 months may be granted for the reduction of IgE-mediated food allergy reactions when all of the following criteria are met:

1. Member is 1 year of age or older.
2. The diagnosis of IgE-mediated food allergy has been confirmed by either of the following:
 - i. Pre-treatment allergen-specific IgE level greater than or equal to 6 IU/mL.
 - ii. Skin-prick test (SPC) with wheal diameter greater than or equal to 4 mm.
3. Member has one of the following:
 - i. A positive physician controlled oral food challenge (e.g., moderate to severe skin, respiratory, or gastrointestinal [GI] symptoms).

- ii. History of a systemic reaction to a food.
- 4. Member has a pre-treatment serum IgE level greater than or equal to 30 IU/mL.
- 5. Member will continue to follow a food-allergen avoidance diet.

D. Chronic spontaneous urticaria

Authorization of 6 months may be granted for treatment of chronic spontaneous urticaria when all of the following criteria are met:

- 1. Member is 12 years of age or older.
- 2. Member remains symptomatic despite treatment with up-dosing (in accordance with EAACI/GA²LEN/EDF/WAO guidelines) of a second-generation H₁ antihistamine (e.g., cetirizine, fexofenadine, levocetirizine, loratadine) for at least 2 weeks.
- 3. Member has been evaluated for other causes of urticaria, including bradykinin-related angioedema and interleukin-1-associated urticarial syndromes (auto-inflammatory disorders, urticarial vasculitis).
- 4. Member has experienced a spontaneous onset of wheals (hives), angioedema, or both, for at least 6 weeks.

E. Immune checkpoint inhibitor-related toxicity

Authorization of 1 month may be granted for treatment of immune checkpoint inhibitor-related toxicity when both of the following are met:

- 1. The member has a refractory case of immune-therapy related severe (G3) pruritus.
- 2. The member has elevated IgE levels.

F. Systemic mastocytosis

Authorization of 12 months may be granted for the treatment of systemic mastocytosis when both of the following are met:

- 1. The major and at least one minor diagnostic criterion for systemic mastocytosis are present or three or more minor diagnostic criteria are present (see Appendix).
- 2. The requested medication will be used in any of the following treatment settings:
 - i. Used as stepwise prophylactic treatment for chronic mast cell mediator-related cardiovascular and pulmonary symptoms when the member has tried both of the following:
 - a. H1 blockers and H2 blockers.
 - b. Corticosteroids.
 - ii. Used for prevention of recurrent unprovoked anaphylaxis.
 - iii. Used for prevention of hymenoptera or food-induced anaphylaxis, with negative specific IgE or negative skin test.
 - iv. Used to improve tolerability of venom immunotherapy.

V. CONTINUATION OF THERAPY

A. Asthma

Authorization of 12 months may be granted for continuation of treatment of asthma when all of the following criteria are met:

- 1. Member is 6 years of age or older.
- 2. Asthma control has improved on Xolair treatment as demonstrated by at least one of the following:
 - i. A reduction in the frequency and/or severity of symptoms and exacerbations.
 - ii. A reduction in the daily maintenance oral corticosteroid dose.
- 3. Member will continue to use maintenance asthma treatments (e.g., inhaled corticosteroid, additional controller) in combination with Xolair.

B. Chronic rhinosinusitis with nasal polyps (CRSwNP)

Authorization of 12 months may be granted for continuation of treatment of CRSwNP when all of the following criteria are met:

1. Member is 18 years of age or older.
2. Member has experienced a positive response as evidenced by improvement in signs and symptoms (e.g., improvement in nasal congestion, nasal polyp size, loss of smell, anterior or posterior rhinorrhea, sinonasal inflammation, hyposmia and/or facial pressure or pain or reduction in corticosteroid use).
3. Member will continue to use a daily intranasal corticosteroid while being treated with the requested medication, unless contraindicated or not tolerated.

C. IgE-mediated food allergy

Authorization of 12 months may be granted for the reduction of IgE-mediated food allergy reactions when all of the following criteria are met:

1. Member is 1 year of age or older.
2. Member has achieved or maintained a positive clinical response to therapy as evidenced by a decrease in hypersensitivity (e.g., moderate to severe skin, respiratory or GI symptoms) to food-allergen.
3. Member will continue to maintain a food-allergen avoidance diet.

D. Chronic spontaneous urticaria

Authorization of 12 months may be granted for continuation of treatment of chronic spontaneous urticaria when all of the following criteria are met:

1. Member is 12 years of age or older.
2. Member has experienced a response (e.g., improved symptoms, decrease in weekly urticaria activity score [UAS7]) since initiation of therapy.

E. Immune checkpoint inhibitor-related toxicities and systemic mastocytosis

All members (including new members) requesting authorization for continuation of therapy must meet all initial authorization criteria.

VI. OTHER

For all indications: Member cannot use the requested medication concomitantly with any other biologic drug or targeted synthetic drug for the same indication.

Note: If the member is a current smoker or vaper, they should be counseled on the harmful effects of smoking and vaping on pulmonary conditions and available smoking and vaping cessation options.

VII. APPENDIX

2017 WHO Diagnostic Criteria for Systemic Mastocytosis

- A. Major Criteria: multifocal, dense infiltrates of mast cells (at least 15 mast cells in aggregates) detected in sections of bone marrow and/or other extracutaneous organs
- B. Minor Criteria
 1. In biopsy sections of bone marrow or other extracutaneous organs, greater than 25% of mast cells in the infiltrate are spindle-shaped or have atypical morphology, or greater than 25% of all mast cells in bone marrow aspirate smears are immature or atypical
 2. Detection of an activating point mutation at codon 816 of *KIT* in the bone marrow, blood, or another extracutaneous organ
 3. Mast cells in bone marrow, blood, or other extracutaneous organs express CD25, with or without CD2, in addition to normal mast cell markers
 4. Serum total tryptase persistently greater than 20 ng/mL (unless there is an associated myeloid neoplasm, in which case this parameter is not valid)

VIII. REFERENCES

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