

PRIOR AUTHORIZATION POLICY

- POLICY:** Allergen Immunotherapy – Grass Pollen Sublingual Products Prior Authorization Policy
- Grastek® (Timothy grass pollen allergen extract sublingual tablets – ALK-Abello)
 - Oralair® (Sweet Vernal, Orchard, Perennial Rye, Timothy, and Kentucky Blue Grass mixed pollens allergen extract sublingual tablets – Stallergenes/Greer)

REVIEW DATE: 10/09/2024

OVERVIEW

Grastek and Oralair are grass pollen allergen extracts indicated for **allergic rhinitis**, with or without conjunctivitis, that has been confirmed by a positive skin test or *in vitro* test for pollen-specific immunoglobulin E (IgE) antibodies for Timothy grass or cross reactive grass pollens (Grastek) or any of the five grasses contained in the product (Oralair).^{1,2} These products are indicated in patients 5 through 65 years of age.

Per product labeling, Grastek must be initiated 12 weeks before the expected onset of each grass pollen season and Oralair must be initiated 4 months before the expected onset of each grass pollen season.^{1,2} Both agents must be continued throughout the season.

Clinical Efficacy

Pivotal trials of Grastek and Oralair included patients with grass pollen-induced allergic rhinitis with or without conjunctivitis confirmed by either a positive skin prick test to Timothy grass pollen or positive *in vitro* test.^{1,2}

POLICY STATEMENT

Prior Authorization is recommended for prescription benefit coverage of Grastek and Oralair. All approvals are provided for the duration noted below.

Automation: None.

RECOMMENDED AUTHORIZATION CRITERIA

Coverage of Grastek and Oralair is recommended in those who meet the following criteria:

FDA-Approved Indication

- 1. Grass Pollen-Induced Allergic Rhinitis.** Approve for 1 year if the patient meets ALL of the following (A, B and C):
 - A)** Patient is ≥ 5 years of age; AND
 - B)** The timing of prescribing meets ONE of the following (i or ii):
 - i.** Grastek: Therapy is initiated 12 weeks prior to the expected onset of the grass pollen season or therapy is being dosed daily continuously for consecutive grass pollen seasons; OR
 - ii.** Oralair: Therapy is initiated 4 months prior to the expected onset of the grass pollen season; AND
 - C)** The diagnosis of grass pollen-induced allergic rhinitis is confirmed by meeting ONE of the following (i or ii):

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- i. Patient has a positive skin test response to a grass pollen from the Pooideae subfamily of grasses (this includes, but is not limited to: sweet vernal, Kentucky blue grass, Timothy grass, orchard, or perennial rye grass); OR
- ii. Patient has a positive *in vitro* test (i.e., a blood test) for allergen-specific immunoglobulin E antibodies for a grass in the Pooideae subfamily of grasses (see examples above).

CONDITIONS NOT RECOMMENDED FOR APPROVAL

Coverage of Grastek and Oralair is not recommended in the following situations:

1. **Concurrent Use of Grastek or Oralair with Subcutaneous Allergen Immunotherapy or Sublingual Allergen Immunotherapy.** Note: This includes allergy shots as well as Odactra[®] [house dust mite {*Dermatophagoides farina* and *Dermatophagoides pteronyssinus*} allergen extract sublingual tablets], Ragwitek[®] [short ragweed pollen allergen extract sublingual tablets]). The efficacy of Grastek and Oralair has not been evaluated in patients who are receiving concomitant allergen immunotherapy.¹ Approved product labeling for both Grastek and Oralair states that concomitant dosing with other allergen immunotherapy may increase the risk of local or systemic adverse events to either subcutaneous or sublingual allergen immunotherapy.
2. Coverage is not recommended for circumstances not listed in the Recommended Authorization Criteria. Criteria will be updated as new published data are available.

REFERENCES

1. Grastek[®] sublingual tablets [prescribing information]. Swindon, Wiltshire, United Kingdom: ALK-Abello A/S; September 2022.
2. Oralair[®] sublingual tablets [prescribing information]. Lenoir, NC: Greer; November 2023.