Clinical Policy: Sweet Vernal, Orchard, Perennial Rye, Timothy, and Kentucky Blue Grass Mixed Pollens Allergen Extract (Oralair)

Reference Number: CP.PMN.85
Effective Date: 11.16.16
Last Review Date: 08.20
Line of Business: Commercial, Medicaid

See Important Reminder at the end of this policy for important regulatory and legal information.

Description
Sweet vernal, orchard, perennial rye, timothy, and Kentucky blue grass mixed pollens allergen extract (Oralair®) is a mixed allergen extract.

FDA Approved Indication(s)
Oralair is indicated as immunotherapy for the treatment of grass pollen-induced allergic rhinitis with or without conjunctivitis confirmed by positive skin test or in vitro testing for pollen-specific IgE antibodies for any of the five grass species contained in this product. Oralair is approved for use in persons 5 through 65 years of age.

Oralair is not indicated for the immediate relief of allergy symptoms.

Policy/Criteria
Provider must submit documentation (such as office chart notes, lab results or other clinical information) supporting that member has met all approval criteria.

It is the policy of health plans affiliated with Centene Corporation® that Oralair is medically necessary when the following criteria are met:

I. Initial Approval Criteria
   A. Allergic Rhinitis (must meet all):
      1. Diagnosis of grass pollen-induced allergic rhinitis;
      2. Prescribed by or in consultation with an allergist or immunologist;
      3. Age ≥ 5 years and ≤ 65 years;
      4. Confirmation of a positive skin test or in vitro testing for pollen-specific IgE antibodies for any of the following grass species:
         a. Sweet vernal;
         b. Orchard;
         c. Perennial rye;
         d. Timothy;
         e. Kentucky blue grass;
      5. Failure of one intranasal corticosteroid, unless clinically significant adverse effects are experienced or all are contraindicated;
      6. Failure of one oral antihistamine at up to maximally indicated doses, unless clinically significant adverse effects are experienced or all are contraindicated;
      7. Dose does not exceed 1 tablet per day.
Approval duration:
Mercy – 12 months
Commercial – Length of Benefit

B. Other diagnoses/indications
1. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): CP.CPA.09 for commercial and CP.PMN.53 for Mercy.

II. Continued Therapy
A. Allergic Rhinitis (must meet all):
1. Currently receiving medication via Centene benefit or member has previously met initial approval criteria;
2. Member is responding positively to therapy;
3. If request is for a dose increase, new dose does not exceed 1 tablet per day.

Approval duration:
Mercy – 12 months
Commercial – Length of Benefit

B. Other diagnoses/indications (must meet 1 or 2):
1. Currently receiving medication via Centene benefit and documentation supports positive response to therapy.
   Approval duration: Duration of request or 12 months (whichever is less); or
2. Refer to the off-label use policy for the relevant line of business if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized): CP.CPA.09 for commercial and CP.PMN.53 for Mercy.

III. Diagnoses/Indications for which coverage is NOT authorized:
A. Non-FDA approved indications, which are not addressed in this policy, unless there is sufficient documentation of efficacy and safety according to the off label use policies – CP.CPA.09 for commercial and CP.PMN.53 for Mercy or evidence of coverage documents.

IV. Appendices/General Information
Appendix A: Abbreviation/Acronym Key
FDA: Food and Drug Administration
IR: index of reactivity

Appendix B: Therapeutic Alternatives
This table provides a listing of preferred alternative therapy recommended in the approval criteria. The drugs listed here may not be a formulary agent for all relevant lines of business and may require prior authorization.
<table>
<thead>
<tr>
<th>Drug Name</th>
<th>Dosing Regimen</th>
<th>Dose Limit/ Maximum Dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>OTC loratadine (Claritin®)</td>
<td>Age 2 to 5 years: 5 mg PO QD Age ≥ 6 years: 10 mg PO QD</td>
<td>10 mg/day</td>
</tr>
<tr>
<td>OTC loratadine-D (Claritin-D® 12 and 24 hour)</td>
<td>Age ≥ 12 years: 1 tablet PO BID (12 hr) QD (24 hr)</td>
<td>10 mg/day</td>
</tr>
<tr>
<td>OTC cetirizine (Zyrtec®)</td>
<td>Age 2 to 5 years: 2.5-5 mg PO QD Age ≥ 6 years: 10 mg PO QD</td>
<td>10 mg/day</td>
</tr>
<tr>
<td>OTC fexofenadine (Allegra Allergy®)</td>
<td>Age 6-months to 2 years: 15 mg PO QD Age 2 to 11 years: 30 mg PO QD Age ≥ 12 years: 60 mg PO BID or 180 mg PO QD</td>
<td>180 mg/day</td>
</tr>
<tr>
<td>fluticasone propionate (Flonase®)</td>
<td>Age ≥ 4 years: 1-2 sprays each nostril QD Age ≥ 12 years: 1-2 sprays each nostril QD</td>
<td>2 sprays each nostril/day</td>
</tr>
<tr>
<td>triamcinolone acetonide (Nasacort AQ®)</td>
<td>Age 2 to 11 years: 1 spray each nostril QD Age ≥ 12 years: 1-2 sprays each nostril QD</td>
<td>Age 2 to 11 years: 1 spray each nostril/day Age ≥ 12 years: 2 sprays each nostril/day</td>
</tr>
<tr>
<td>mometasone furoate monohydrate (Nasonex®)</td>
<td>Age 2 to 11 years: 1 spray each nostril QD Age ≥ 12 years: 2 sprays each nostril QD</td>
<td>Age 2 to 11 years: 1 spray each nostril/day Age ≥ 12 years: 2 sprays each nostril/day</td>
</tr>
</tbody>
</table>

*Therapeutic alternatives are listed as Brand name® (generic) when the drug is available by brand name only and generic (Brand name®) when the drug is available by both brand and generic.*

**Appendix C: Contraindications/Boxed Warnings**
- Contraindication(s): severe, unstable or uncontrolled asthma; history of eosinophilic esophagitis; history of any severe systemic allergic reaction or any severe local reaction to sublingual allergen immunotherapy; hypersensitivity to any of the inactive ingredients contained in this product
- Boxed warning(s): severe allergic reactions

**V. Dosage and Administration**

<table>
<thead>
<tr>
<th>Indication</th>
<th>Dosing Regimen</th>
<th>Maximum Dose</th>
</tr>
</thead>
<tbody>
<tr>
<td>Grass pollen-induced allergic rhinitis</td>
<td>Age 5 to 17 years: 100 IR (index of reactivity) sublingually (SL) on day 1 followed by 200 IR SL on day 2 and 300 IR SL QD on day 3 and thereafter.</td>
<td>300 IR/day</td>
</tr>
</tbody>
</table>

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### Indication

Age 18 to 65 years: 300 IR (index of reactivity) SL QD

Treatment should be initiated 4 months before the expected onset of each grass pollen season and continue treatment throughout the season

### VI. Product Availability

Tablets: 100 IR, 300 IR

### VII. References


### Reviews, Revisions, and Approvals

<table>
<thead>
<tr>
<th>Date</th>
<th>P&amp;T Approval Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>01.12.17</td>
<td>11.17</td>
</tr>
<tr>
<td>04.02.18</td>
<td>08.18</td>
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</table>

Converted to new template. Minor changes to verbiage and grammar. References updated.

3Q 2018 annual review: Medicaid and Commercial (CP.CPA.111) policies combined; age added to policy; Medicaid: increased initial approval duration to 12 months; Commercial: removed leukotriene
Clinical Policy
Sweet Vernal, Orchard, Perennial Rye, Timothy, and Kentucky Blue Grass Mixed Pollens Allergen Extract

Reviews, Revisions, and Approvals

<table>
<thead>
<tr>
<th>Description</th>
<th>Date</th>
<th>P&amp;T Approval Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>modifiers as pdl alternative per 2017 guidelines; references reviewed and updated.</td>
<td></td>
<td></td>
</tr>
<tr>
<td>No significant changes: pediatric age requirement expanded down to age 5 years for 10 years old per drug labeling changes; references reviewed and updated.</td>
<td>12.19.18</td>
<td>08.19</td>
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<tr>
<td>3Q 2019 annual review: no significant changes; corrected age restriction from &lt; 65 years to ≤ 65 years per PI; references reviewed and updated.</td>
<td>04.22.19</td>
<td>08.19</td>
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<tr>
<td>3Q 2020 annual review: no significant changes; references reviewed and updated.</td>
<td>04.06.20</td>
<td>08.20</td>
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</table>

Important Reminder
This clinical policy has been developed by appropriately experienced and licensed health care professionals based on a review and consideration of currently available generally accepted standards of medical practice; peer-reviewed medical literature; government agency/program approval status; evidence-based guidelines and positions of leading national health professional organizations; views of physicians practicing in relevant clinical areas affected by this clinical policy; and other available clinical information. The Health Plan makes no representations and accepts no liability with respect to the content of any external information used or relied upon in developing this clinical policy. This clinical policy is consistent with standards of medical practice current at the time that this clinical policy was approved. “Health Plan” means a health plan that has adopted this clinical policy and that is operated or administered, in whole or in part, by Centene Management Company, LLC, or any of such health plan’s affiliates, as applicable.

The purpose of this clinical policy is to provide a guide to medical necessity, which is a component of the guidelines used to assist in making coverage decisions and administering benefits. It does not constitute a contract or guarantee regarding payment or results. Coverage decisions and the administration of benefits are subject to all terms, conditions, exclusions and limitations of the coverage documents (e.g., evidence of coverage, certificate of coverage, policy, contract of insurance, etc.), as well as to state and federal requirements and applicable Health Plan-level administrative policies and procedures.

This clinical policy is effective as of the date determined by the Health Plan. The date of posting may not be the effective date of this clinical policy. This clinical policy may be subject to applicable legal and regulatory requirements relating to provider notification. If there is a discrepancy between the effective date of this clinical policy and any applicable legal or regulatory requirement, the requirements of law and regulation shall govern. The Health Plan retains the right to change, amend or withdraw this clinical policy, and additional clinical policies may be developed and adopted as needed, at any time.

This clinical policy does not constitute medical advice, medical treatment or medical care. It is not intended to dictate to providers how to practice medicine. Providers are expected to exercise
professional medical judgment in providing the most appropriate care, and are solely responsible for the medical advice and treatment of members. This clinical policy is not intended to recommend treatment for members. Members should consult with their treating physician in connection with diagnosis and treatment decisions.

Providers referred to in this clinical policy are independent contractors who exercise independent judgment and over whom the Health Plan has no control or right of control. Providers are not agents or employees of the Health Plan.

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Note:
For Medicaid members, when state Medicaid coverage provisions conflict with the coverage provisions in this clinical policy, state Medicaid coverage provisions take precedence. Please refer to the state Medicaid manual for any coverage provisions pertaining to this clinical policy.

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