Clinical Policy: IncobotulinumtoxinA (Xeomin)
Reference Number: ERX.SPA.194
Effective Date: 01.11.17
Last Review Date: 05.18

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

Description
IncobotulinumtoxinA (Xeomin®) is an acetylcholine release inhibitor and a neuromuscular blocking agent.

FDA Approved Indication(s)
Xeomin is indicated for the treatment or improvement of adult patients with:
- Upper limb spasticity
- Cervical dystonia (CD) in both botulinum toxin-naïve and previously treated patients
- Blepharospasm in adults previously treated with onabotulinumtoxinA (Botox®)
- Temporary improvement in the appearance of moderate to severe glabellar lines associated with corrugator and/or procerus muscle activity in adult patients

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 Envolve Pharmacy Solutions™ that Xeomin is medically necessary when the following criteria are met:

I. Initial Approval Criteria
   A. Cervical Dystonia (must meet all):
      1. Diagnosis of CD (see Appendix C);
      2. Prescribed by or in consultation with a neurologist, orthopedist, physiatrist, or physical medicine and rehabilitation specialist;
      3. Age ≥ 18 years;
      4. Experiencing involuntary contractions of the neck and shoulder muscles (e.g., splenius, sternocleidomastoid, levator scapulae, scalene, trapezius, semispinalis capitis) resulting in abnormal postures or movements of the neck, shoulder or head;
      5. Contractions are causing pain and functional impairment;
      6. Provider submits treatment plan detailing the quantity (in units) of Xeomin to be injected in each muscle site, anticipated frequency of injection, and total dose per visit;
      7. Dose does not exceed 120 units per treatment session.
      Approval duration: 12 weeks (single treatment session)

   B. Blepharospasm (a focal dystonia) (must meet all):
      1. Diagnosis of blepharospasm (i.e., abnormal contraction of eyelid muscles);
      2. Prescribed by or in consultation with a neurologist or ophthalmologist;
      3. Age ≥ 18 years;
      4. Member previously received treatment with onabotulinumtoxinA (Botox);
      5. Provider submits treatment plan detailing the quantity (in units) of Xeomin to be injected in each muscle site, anticipated frequency of injection, and total dose per visit;
      6. Dose does not exceed 35 units per eye per treatment session.
      Approval duration: 12 weeks (single treatment session)

   C. Upper Limb Spasticity (must meet all):
      1. Diagnosis of upper limb spasticity;
      2. Prescribed by or in consultation with a neurologist, orthopedist, physiatrist, or physical medicine and rehabilitation specialist;
      3. Age ≥ 18 years;
4. Provider submits treatment plan detailing the quantity (in units) of Xeomin to be injected in each muscle site, anticipated frequency of injection, and total dose per visit;
5. Dose does not exceed 400 units per treatment session.
Approval duration: 12 weeks (single treatment session)

D. Other diagnoses/indications
1. Refer to ERX.PA.01 if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized).

II. Continued Therapy
A. All Indications in Section I (must meet all):
   1. Currently receiving medication via a health plan affiliated with Envolve Pharmacy Solutions or member has previously met initial approval criteria;
   2. Member is responding positively to therapy;
   3. It has been at least 12 weeks since the last injection of Xeomin;
   4. Provider submits treatment plan detailing the quantity (in units) of Xeomin to be injected in each muscle site, anticipated frequency of injection, and total dose per visit;
   5. If request is for a dose increase, new dose does not exceed the following indication-specific maximums (a or b):
      a. CD: 120 units per treatment session;
      b. Upper limb spasticity: 400 units per treatment session;
      c. Blepharospasm: 35 units per eye per treatment session.
Approval duration: 12 weeks (single treatment session)

B. Other diagnoses/indications (must meet 1 or 2):
   1. Currently receiving medication via a health plan affiliated with Envolve Pharmacy Solutions and documentation supports positive response to therapy.
      Approval duration: 12 weeks (single treatment session); or
   2. Refer to ERX.PA.01 if diagnosis is NOT specifically listed under section III (Diagnoses/Indications for which coverage is NOT authorized).

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 policy – ERX.PA.01 or evidence of coverage documents;
B. Cosmetic treatment of hyperfunctional wrinkles of the upper face including glabellar frown lines, deep forehead wrinkles and periorbital wrinkles (crow’s feet).

IV. Appendices/General Information
Appendix A: Abbreviation/Acronym Key
CD: cervical dystonia
FDA: Food and Drug Administration

Appendix B: Therapeutic Alternatives
Not applicable

Appendix C: Definition and Classification of Dystonia
Dystonia is defined as a movement disorder characterized by sustained or intermittent muscle contractions causing abnormal, often repetitive, movements, postures, or both.
- Dystonic movements are typically patterned and twisting, and may be tremulous.
- Dystonia is often initiated or worsened by voluntary action and associated with overflow muscle activation.

Dystonia is classified along two axes:
CLINICAL POLICY
IncobotulinumtoxinA

- Clinical characteristics: Age at onset, body distribution, temporal pattern, associated features (additional movement disorders or neurological features) - the clinical characteristics fall into several specific dystonia syndromes that help to guide diagnosis and treatment;
- Etiology: Nervous system pathology, inheritance.

V. Dosage and Administration

<table>
<thead>
<tr>
<th>Indication</th>
<th>Dosing Regimen</th>
<th>Maximum Dose</th>
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<tbody>
<tr>
<td>CD</td>
<td>The usual starting dose is 120 units per treatment session, doses up to 300</td>
<td>120 units/treatment session</td>
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<td>units may be used in treatment-experienced patients. Dose, number, and location</td>
<td>of muscles involved, severity of dystonia,</td>
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<td></td>
<td>of injection sites should be based on the number and location of muscles</td>
<td>and response to any previous botulinum</td>
</tr>
<tr>
<td></td>
<td>involved, severity of dystonia, and response to any previous botulinum</td>
<td>toxin injections.</td>
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<td>toxin injections.</td>
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<tr>
<td>Blepharospasm</td>
<td>When initiating Xeomin therapy, the dose, number, and location of injections</td>
<td>35 units/eye/treatment session</td>
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<td>should be based on the previous dosing of Botox. If the previous dose of</td>
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<td></td>
<td>Botox is not known, the recommended starting dose is 1.25-2.5 units per</td>
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<td>injection site.</td>
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<td>Upper limb spasticity</td>
<td>Dosing varies based on location of muscles to be treated</td>
<td>400 units/treatment session</td>
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<td>(refer to dosing chart in the prescribing information).</td>
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VI. Product Availability
Vials: 50 units, 100 units, 200 units

VII. References

Reviews, Revisions, and Approvals

<table>
<thead>
<tr>
<th>Policy created</th>
<th>Date</th>
<th>P&amp;T Approval Date</th>
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<tbody>
<tr>
<td>12.01.16</td>
<td>01.17</td>
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<tr>
<td>4Q17 Annual Review</td>
<td>10.03.17</td>
<td>11.17</td>
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<tr>
<td>- Converted to new template.</td>
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<tr>
<td>- Added requirement for documentation of positive response to therapy, for reauthorization.</td>
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<tr>
<td>2Q 2018 annual review: added physical medicine and rehabilitation specialist for relevant indications; broke up cervical dystonia and upper limb spasticity into separate criteria sets; required prescriber submission of treatment plan details for all initial and continued approval indications; references reviewed and updated.</td>
<td>02.09.18</td>
<td>05.18</td>
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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.

This Clinical Policy is not intended to dictate to providers how to practice medicine, nor does it constitute a contract or guarantee regarding payment or results. 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.

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