

POLICY NAME	Botox (onabotulinumtoxin A)	POLICY #	2373P
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Criteria

Criteria for Coverage for Chronic Migraine Headaches

- ☐ **1.1** Documented diagnosis of chronic migraine.
- ☐ **1.2** Documented headache diary or chart notes describing the patient's migraine history.
- ☐ **1.3** Documented failure, intolerance, or contraindication to at least 2 American Headache Society Level A or B migraine prophylactic therapies with claims history to support member compliance with filling at least a 90 day supply within a 120 day time frame
 - Beta Blockers [?] Level A: metoprolol, propranolol, timolol [?] Level B: atenolol, nadolol
 - Antidepressants [?] Level B: amitriptyline, nortriptyline, duloxetine, venlafaxine
 - Anticonvulsants [?] Level A: divalproex, valproic acid, topiramate
- ☐ **1.4** Reauthorization requires a documented reduction in “migraine days” by 7 days per month
- ☐ **1.5** Prescribed by a neurologist (central nervous system doctor), physical medicine rehabilitation specialist, or pain management specialist
- ☐ **1.6** Approval Time
 - Initial approval: 4 procedures each spaced 12 weeks apart within a 12 month approval duration
 - Subsequent approvals: 4 procedures, spaced apart by 12 weeks with documentation that patient has experienced a positive response to therapy [?] Reduction in headache frequency and/or intensity [?] Use of acute migraine medications (e.g., non-steroidal anti-inflammatory drugs (NSAIDs), triptans) has decreased since the start of Botox therapy [?] Documentation that patient continues to be monitored for medication overuse headache

Coverage Criteria for Concurrent use of a Prophylactic C-GRP and Botulinum toxin

- ☐ **2.1** Documentation showing that member has had at least a 6 month trial of botulinum toxin without adequate improvement in migraine, OR
- ☐ **2.2** Documentation showing that member has had at least a 3 month trial of Aimovig, Ajovy, Emgality, Nurtec, Qulipta, or Vyepti as prophylactic treatment without adequate improvement in migraine
 - Coverage of Emgality 120mg requires trial and failure of Aimovig and Ajovy

Criteria for Coverage for Cervical Dystonia

- ☐ **3.1** Alternative diagnoses ruled out including adverse effects of medications or other injuries or disorders of the muscles, nerves, tendons, joints, cartilage, or spinal discs Criteria
- ☐ **3.2** Involuntary contractions of the neck muscles
- ☐ **3.3** Chronic head torsion (twisting) or tilt
- ☐ **3.4** Symptoms present for at least 6 months
- ☐ **3.5** Approval Time
 - Initial Approval: 4 procedures each spaced 12 weeks apart within a 12 month approval duration
 - Reapproval: 4 procedures each spaced 12 weeks apart with documentation that patient experienced a positive response to therapy

Criteria for Coverage for Overactive Bladder Syndrome

- ☐ **4.1** Documented urinary urgency and frequency, urge incontinence and/or waking up in the night to urinate;
- ☐ **4.2** Documented limited ability to participate in daily activities
- ☐ **4.3** Documented failure of conservative therapies
 - Pelvic floor exercises
 - Biofeedback
 - Times voids
 - Dietary/fluid management under the direction of a qualified therapist
- ☐ **4.4** Prescribed by a urologist (urinary tract doctor)
- ☐ **4.5** Documented failure, intolerance, or contraindication to at least 2 anticholinergics, OR
 - Some examples are oxybutynin, tolterodine, Enablex, Toviaz
- ☐ **4.6** Documented failure, intolerance, or contraindication to 1 anticholinergic and 1 other class of medication for overactive bladder syndrome
 - Some examples are amitriptyline, desipramine, clonidine, Myrbetriq, duloxetine
- ☐ **4.7** Approval Time
 - Initial Approval: 4 procedures each spaced 12 weeks apart within a 12 month approval duration
 - Reapproval: 4 procedures each spaced 12 weeks apart with documentation that patient experienced a positive response to therapy

Criteria for Coverage for Dynamic Contracture in Cerebral Palsy

- ☐ **5.1** Documented hygienic problems or significant functional limitations
- ☐

5.2 Approval Time

- Initial Approval: 4 procedures each spaced 12 weeks apart within a 12 month approval duration
- Reapproval: 4 procedures each spaced 12 weeks apart

Criteria for Coverage for Axillary Hyperhidrosis (excessive perspiration of the underarms)

- ☐ 6.1 Uncontrolled perspiration present for more than 1 year
- ☐ 6.2 Perspiration severely impacts the member's occupational and social activities
- ☐ 6.3 Documented failure, intolerance, or contraindication to an adequate trial of topical aluminum chloride solution
- ☐ 6.4 Documented failure, intolerance, or contraindication to local and systemic drug therapy
 - Anticholinergics
 - Beta blockers
 - Benzodiazepines
- ☐ 6.5 Botox is not covered for hyperhidrosis (excessive perspiration) in other body areas because safety and efficacy has not been established
- ☐ 6.6 Approval Time
 - Initial Approval: 4 procedures each spaced 12 weeks apart within a 12 month approval duration
 - Reapproval: 4 procedures each spaced 12 weeks apart with documentation that patient experienced a positive response to therapy's

Criteria for Coverage for Chronic Anal Fissures

- ☐ 7.1 Documented trial and failure of conservative therapy
 - Nitroglycerin ointment
 - Diltiazem
 - Bethanechol
- ☐ 7.2 Prescribed by a gastroenterologist (stomach doctor) or colorectal (colon and anus) surgeon;
- ☐ 7.3 Approval Time
 - Initial Approval: 2 procedures spaced 12 weeks apart within a 12 month approval duration
 - Max 2 procedures per lifetime

Criteria for Coverage for Upper Limb Spasticity

- ☐ 8.1 Documented focal wrist, elbow, or finger spasticity which originated at least 6 weeks post-cerebrovascular event (CVE) or progression of multiple sclerosis
- ☐ 8.2 Difficulty maintaining hygiene, dressing or pain

- ☐ **8.3 Documented failure, intolerance, or contraindication to oral antispasmodics and muscle relaxants**
 - Baclofen
 - Tizanidine
 - Cyclobenzaprine
 - Methocarbamol
 - Carisoprodol
- ☐ **8.4 Sufficient motivation and cognitive function to actively participate in physical therapy post injection**
- ☐ **8.5 No documented fixed contractures (tightening of muscle tendons, ligaments or skin which prevents normal movement of the body part) or profound muscle wasting; AND**
- ☐ **8.6 Member will not receive treatment with phenol, alcohol, or surgery**
- ☐ **8.7 Approval Time**
 - Initial Approval: 4 procedures each spaced 12 weeks apart within a 12 month approval duration
 - Reapproval: 4 procedures each spaced 12 weeks apart with documentation that patient experienced a positive response to therapy

Criteria for Coverage of Upper or Lower Limb Spasticity for Pediatric Patients

- ☐ **9.1 Age 2 to 17 years**
- ☐ **9.2 Documented upper limb spasticity due to cerebral palsy, traumatic brain injury, multiple sclerosis, spinal cord injury, and stroke**
- ☐ **9.3 Approval time**
 - Initial Approval: 4 procedures each spaced 12 weeks apart within a 12 month approval duration
 - Reapproval: 4 procedures each spaced 12 weeks apart with documentation that patient experienced a positive response to therapy

Criteria for Coverage for Lower Limb Spasticity

- ☐ **10.1 Documented severe spastic equinovarus foot (overactivity of lower leg muscles) as a result of stroke**
- ☐ **10.2 Failure to respond to oral antispasmodics, physical therapy, orthotics or other non-operative modalities**
 - Some examples of antispasmodics are baclofen, tizanidine, cyclobenzaprine
- ☐ **10.3 Sufficient motivation and cognitive function to actively participate in physical therapy post injection**
- ☐ **10.4 No documented fixed contractures or profound muscle atrophy**
- ☐ **10.5 Member will not receive treatment with phenol, alcohol, or surgery**

☐ **10.6 Approval Time**

☐ **10.7 Initial Approval:** 4 procedures each spaced 12 weeks apart within a 12 month approval duration

- Reapproval: 4 procedures each spaced 12 weeks apart with documentation that patient experienced a positive response to therapy

Criteria for Coverage for Writer's Cramp (abnormal movement of the hand and/or forearm during tasks requiring skilled hand use, such as writing)

☐ **11.1 Documented significant functional limitations that interfere with daily activities**

☐ **11.2 Documented failure of conservative treatments;**

- Transcutaneous electrical nerve stimulation
 - Biofeedback References
 - Hypnotherapy
 - Relaxation therapy
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☐ **11.3 Approval Time**

- Initial Approval: 4 procedures each spaced 12 weeks apart within a 12 month approval duration
- Reapproval: 4 procedures each spaced 12 weeks apart with documentation that patient experienced a positive response to therapy

Criteria for Coverage of Pediatric Detrusor Overactivity associated with a Neurologic Condition

☐ **12.1 Age 5 and older**

☐ **12.2 Documented inadequate response to or intolerance of anticholinergic medications**

- Initial Approval: 4 procedures each spaced 12 weeks apart within a 12 month approval duration
- Reapproval: 4 procedures each spaced at least 12 weeks apart with documentation of positive response to therapy

Criteria for Coverage for Other Indications

☐ **13.1 Diagnosis of Achalasia**

- muscle disorder which prevents lower esophagus to open up during swallowing
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☐ **13.2 Diagnosis of Adductor laryngeal dystonia**

- abnormal involuntary excessive contraction of the muscles that bring the vocal cords together
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☐ **13.3 Diagnosis of Blepharospasm**

- abnormal contraction of the eyelid muscles
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☐

13.4 Diagnosis of Focal dystonia

- Neuromuscular disorder with involuntary muscle contractions in one body part such as neck, face, jaw, feet or hands

☐ 13.5 Diagnosis of Hemifacial spasm

- neuromuscular disorder causing frequent involuntary contractions of the muscles on one side of the face

☐ 13.6 Diagnosis of Jaw closing dystonia

- involuntary and forceful muscle contractions of the face, jaw, and/or tongue

☐ 13.7 Diagnosis of Strabismus

- condition in which the eyes do not properly align with each other when looking at an object

☐ 13.8 Approval Time

- Initial Approval: 4 procedures each spaced 12 weeks apart within a 12 month approval duration
- Reapproval: 4 procedures each spaced 12 weeks apart with documentation that patient experienced a positive response to therapy CPT Codes HCPCS Codes J0585 Injection, onabotulinumtoxinA [Botox]