

NC Pharmacy Prior Approval Request for Neuromuscular Blocking Agents: Botox/Myobloc/Dysport/Xeomin

Beneficiary Information

1. Beneficiary Last Name: _____ 2. First Name: _____
3. Beneficiary ID #: _____ 4. Beneficiary Date of Birth: _____ 5. Beneficiary Gender: _____

Prescriber Information

6. Prescribing Provider NPI #: _____ Provider Fax #: _____
7. Requester Contact Information - Name: _____ Phone #: _____ Ext. _____

Drug Information

8. Drug Name: _____ 9. Strength: _____ 10. Quantity Per 30 Days: _____
11. Length of Therapy (in days): up to 30 Days 60 Days 90 Days 120 Days 180 Days 365 Days

Clinical Information

1. What is the prescribed dosage? _____ units per _____ days
 2. What is the diagnosis or indication for the medication?
 - Blepharospasm (**Botox, Dysport, Xeomin**)
 - Disorders of eye movement (strabismus) (**Botox**)
 - Spasmodic torticollis, secondary to cervical dystonia (**Botox, Dysport, Myobloc, Xeomin**)
 - Spasticity in beneficiaries age 2 and up (**Botox**)
 - Severe axillary hyperhidrosis (ANSWER QUESTIONS 2 AND 3 BELOW) (**Botox, Dysport**)
 - Sialorrhea (**Botox, Myobloc**)
 - Chronic Sialorrhea in beneficiaries age 2 and up (**Xeomin**)
 - Chronic anal fissure refractory to conservative treatment (**Botox**)
 - Esophageal achalasia recipients in whom surgical treatment is not indicated (**Botox**)
 - Infantile cerebral palsy, specified or unspecified (**Botox**)
 - Hemifacial Spasms (**Botox, Dysport**)
 - Laryngeal dystonia and adductor spasmodic dysphonia (**Botox**)
 - Upper limb spasticity in adults (**Dysport, Xeomin**)
 - Upper limb spasticity in pediatric beneficiaries 2 years of age and older, excluding spasticity caused by cerebral palsy (**Dysport**)
 - Lower limb spasticity in adults and pediatric beneficiaries 2 years of age and older (**Dysport**)
 - Upper limb spasticity in pediatric beneficiaries 2 to 17 years of age, excluding spasticity caused by cerebral (**Xeomin**)
 2. Does the patient have documented medical complications due to hyperhidrosis? Yes No Please List: _____
 3. Has the patient failed a 6-month trial of conservative management including the use of topical aluminum chloride or extra strength antiperspirant?
 - Yes No Please List product (s) tried: _____
- Chronic Migraine (18 and older) New Therapy (approval up to 6 months) (BOTOX)**
4. Does the patient have 15 or more days each month with headache lasting 4 or more hours? Yes No
 5. Has the patient tried and failed prophylactic medications from at least 3 different drug classes (beta blockers, calcium channel Blockers, tricyclic antidepressants and anticonvulsants) each for at least 3 months of therapy? Yes No List meds tried: _____
- Chronic Migraine Continuation of Therapy (approval up to 1 year) (BOTOX)**
6. Has the patient responded favorably after the first 2 injections? Yes No
 7. Has the average number of headache days decreased by 6 or more days from the patient's baseline headache frequency? Yes No
- Urinary Incontinence (Botox)**
8. Does the patient have detrusor overactivity associated with neurologic conditions? Yes No
 9. Has the patient tried and failed an anticholinergic medication? Yes No List med tried: _____
 10. Does the patient have a documented contraindication, intolerable side effects, or allergy to anticholinergic medications? Yes No
- Overactive Bladder (BOTOX)**
11. Has the beneficiary tried and failed on 2 anticholinergic medications? Yes No List meds tried _____
 12. Does the beneficiary have a documented contraindication, intolerable side effect, or allergy to anticholinergic medications? Yes No

Signature of Prescriber: _____ Date: _____

(Prescriber Signature Mandatory)

I certify that the information provided is accurate and complete to the best of my knowledge, and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.