

NC Medicaid and NC Health Choice 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 #:		
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
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Clinical Information

 What is the prescribed dosage? units per days 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 3 AND 4 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)				
3. Does the patient have documented medical complications due to hyperhidrosis?				
□ Yes □ No Please List:				
4. 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)				



5. Does the patient have 15 or more days each month with headache lasting 4 or more hours? □ Yes □ No 6. 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)
7. Has the patient responded favorably after the first 2 injections? □ Yes □ No
8. 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)

9. Does the patient have detrusor overactivity associated with neurologic conditions? □ Yes □ No
10. Has the patient tried and failed an anticholinergic medication? □ Yes □ No List meds tried

11. Does the patient have a documented contraindication, intolerable side effects, or allergy to anticholinergic medications?

Yes
No

Overactive Bladder (BOTOX)

12. Has the beneficiary tried and failed on 2 anticholinergic medications?

Yes
No List meds tried

13. 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.

Please fax this form to 1-800-678-3189 Pharmacy PA Call Center: 1-866-799-5318