



Fax Referral To: 800-323-2445
Phone: 866-278-6634

Botulinum Toxin Type A and B Enrollment Form

For Blue Cross Blue Shield of Rhode Island Members

Date: _____

Needs by Date (Please Specify): _____

Ship to: ☐ Patient ☐ Office ☐ Other: _____

PATIENT INFORMATION

(Complete the following or send patient demographic sheet)

Patient Name: _____
Address: _____
City, State, Zip: _____
Home Phone: _____
Alternate Phone: _____
SS #: _____
Insurance ID: _____ Primary Language: _____
Date of Birth: _____ Gender: _____

PRESCRIBER INFORMATION

Prescriber's Name: _____
State License #: _____ UPIN: _____
DEA #: _____ NPI #: _____
Group or Hospital: _____
Address: _____
City, State Zip: _____
Phone: _____ Fax: _____
Contact Person: _____
Contact Phone: _____

INSURANCE INFORMATION (Please copy and attach the front and back of insurance and prescription drug card)

Prescription Card:	Name of Insurer: _____	ID#: _____	BIN: _____	PCN: _____	Group: _____
Primary Insurance:	Subscriber: _____	ID#: _____	Name of Insurer: <u>Blue Cross Blue Shield of RI</u>	Phone: _____	
Secondary Insurance:	Subscriber: _____	ID#: _____	Name of Insurer: _____	Phone: _____	

STATEMENT OF MEDICAL NECESSITY for BCBS of Rhode Island Members

Diagnosis (ICD-9 Code and description): ☐ _____ • Date of Diagnosis: _____

APPROVAL CRITERIA: CHECK ALL BOXES THAT APPLY.

NOTE: Any areas not filled out are considered not applicable to your patient & MAY AFFECT THE OUTCOME of this request.

- Patient has a diagnosis of **Strabismus, Achalasia, or Chronic Anal Fissures**? ☐ Yes ☐ No
- Patient has one of the following disorders associated with spasticity or Dystonia? ☐ Yes ☐ No
 - ☐ Blepharospasm
 - ☐ Hereditary Spastic Paraplegia
 - ☐ Multiple Sclerosis
 - ☐ Schilder's Disease
 - ☐ Spasticity form Stroke or Spinal Cord Injury
 - ☐ Idiopathic Torsion Dystonia
 - ☐ Orofacial Dyskinesia (i.e., jaw closure dystonia)
 - ☐ Symptomatic Torsion Dystonia
 - ☐ Cervical Dystonia
 - ☐ Equinus foot, if related to Cerebral Palsy
 - ☐ Infantile Cerebral Palsy
 - ☐ Neuromyelitis Optica
 - ☐ Spastic Hemiplegia
 - ☐ Upper Limb spasticity
 - ☐ Organic Writer's Cramp
 - ☐ Spasmodic Dysphonia/Laryngeal Dystonia*
 - ☐ Facial Nerve (VII) Dystonia
 - ☐ Forms of Upper Motor Neuron Spasticity
- Patient has drooling (sialorrhea) associated with Parkinson's disease. ☐ Yes ☐ No
- Patient has Cervical Dystonia? (If Yes, please answer #1-6 below) ☐ Yes ☐ No
 - 1 - Prior to the initiation of therapy, was the cervical dystonia moderate in severity or greater? ☐ Yes ☐ No
 - 2 - Does the patient have a history of recurrent clonic and/or tonic involuntary contractions in the sternocleidomastoid, splenius, trapezius, or posterior cervical muscles? ☐ Y ☐ N
 - 3 - Prior to the initiation of therapy, did the patient exhibit sustained head tilt and/or abnormal posturing with limited range of motion in the neck? ☐ Yes ☐ No
 - 4 - Is the duration of cervical dystonia greater than 6 months? ☐ Yes ☐ No
 - 5 - Has the patient previously received botulinum toxin? ☐ Yes ☐ No
 - 6 - If Yes, Did the patient respond to the initial treatment and was this response documented in the medical record? ☐ Yes ☐ No
- Patient has incontinence related to bladder/detrusor overactivity or incontinence of neurogenic origin? ☐ Yes ☐ No
 - Patient has an inadequate response to anticholinergic therapy? ☐ Yes ☐ No
- Patient has bladder/detrusor sphincter dyssynergia of neurogenic origin? ☐ Yes ☐ No
- Patient has hyperhidrosis that is incapacitating or causing persistent chronic cutaneous conditions? ☐ Yes ☐ No
- Patient has had an inadequate response to a 6-month trial of non-surgical treatment for hyperhidrosis? ☐ Yes ☐ No

* A disorder of speech due to abnormal control of the laryngeal muscles present only during the specific task of speaking.

****NOTES:** Botulinum Toxin is **not approvable** for the following, as well as other conditions:

- | | |
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| <ul style="list-style-type: none">• Skin Wrinkles or other cosmetic indications• Headache, except for prevention of chronic migraine• Anismus• Chronic Motortic Disorder• Fibromyalgia/Fibromyositis• Gastroparesis• Low Back Pain• Myofacial Pain Syndrome | <ul style="list-style-type: none">• Neck Pain not related to conditions mentioned above• Parkinson's Disease• Tics associated with Tourette's Syndrome• Tourette's Syndrome• Tremors• Urinary and Anal Spinchter Dysfunction• Stuttering• Carpal Tunnel Syndrome |
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****For chronic migraine, physicians requesting initial therapy or renewals are required to provide information from the headache diary or other evidence of the patient's headache frequency and duration*****

- Does the patient experience headache on ≥ 15 days per month? ☐ Yes ☐ No
- Do headaches last ≥ 4 hours? ☐ Yes ☐ No
- Please document the patient's pre-treatment monthly headache frequency and daily headache duration.
- Has the patient experienced headaches for ≥ 3 months? ☐ Yes ☐ No
- Has the patient been evaluated for medication overuse headache and treated as appropriate? ☐ Yes ☐ No
- Did the patient complete adequate trials (each ≥ 8 weeks) of at least 2 oral preventative agents from different pharmacological classes? ☐ Yes ☐ No
- Please document oral preventative agents tried.
 - a. Divalproex sodium ☐ Yes ☐ No
 - b. Topiramate ☐ Yes ☐ No
 - c. Gabapentin ☐ Yes ☐ No
 - d. Amitriptyline ☐ Yes ☐ No
 - e. Venlafaxine ☐ Yes ☐ No
 - f. Fluoxetine ☐ Yes ☐ No
 - g. Propranolol, timolol, atenolol, metoprolol, or nadolol ☐ Yes ☐ No
 - h. Nimodipine, verapamil or diltiazem ☐ Yes ☐ No
 - i. Lisinopril ☐ Yes ☐ No
 - j. Candesartan ☐ Yes ☐ No
 - k. Other (please document) ☐ Yes ☐ No
- If patient has not tried 2 oral agents, does the patient have a contraindication to each pharmacological class of agents not tried? ☐ Yes ☐ No
- If not contraindicated, is there another clinical reason(s) for not completing adequate trials of all pharmacological classes? ☐ Yes ☐ No
- Please document the agent(s) tried and the reason(s) for discontinuation.
 - a. Allergic reaction ☐ Yes ☐ No
 - b. Adverse event ☐ Yes ☐ No
 - c. Unable to tolerate medication/unpleasant side effect ☐ Yes ☐ No
 - d. Significant drug interaction ☐ Yes ☐ No
 - e. Other (please document below) ☐ Yes ☐ No

- If patient is receiving botulinum toxin therapy, how many days per month has the headache frequency decreased since starting therapy? _____
- If patient is receiving botulinum toxin therapy, how many hours per month has the headache duration decreased since starting therapy? _____

PRESCRIPTION INFORMATION				
MEDICATION	STRENGTH	DIRECTIONS	QUANTITY	REFILLS
<input type="checkbox"/> Botox				
<input type="checkbox"/> Dysport				
<input type="checkbox"/> Myobloc				
<input type="checkbox"/> Xeomin				

X	X
PRODUCT SUBSTITUTION PERMITTED	DISPENSE AS WRITTEN
(Date)	(Date)