

Myobloc

Prior Authorization Request

CVS Caremark administers the prescription benefit plan for the patient identified. This patient's benefit plan requires prior authorization for certain medications in order for the drug to be covered. To make an appropriate determination, providing the most accurate diagnosis for the use of the prescribed medication is necessary. **Please respond below and fax this form to CVS Caremark toll-free at 1-855-330-1720**. If you have questions regarding the prior authorization, please contact CVS Caremark at **1-888-877-0518**. For inquiries or questions related to the patient's eligibility, drug copay or medication delivery; please contact the Specialty Customer Care Team: CaremarkConnect[®] 1-800-237-2767.

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Patient's Name:	Date:
Patient's ID:	
Physician's Name:	
Specialty:	
Physician Office Telephone:	Physician Office Fax:
<u>Referring</u> Provider Info: Same as Requ	esting Provider
Name:	NPI#:
Fax:	Phone:
Rendering Provider Info:	ring Provider 🛛 Same as Requesting Provider
Name:	NPI#:
Fax:	Phone:

Approvals may be subject to dosing limits in accordance with FDA-approved labeling, accepted compendia, and/or evidence-based practice guidelines.

Required Demographic Information:

Patient Weight: _____kg

Patient Height: _____cm

Please indicate the place of service for the requested drug: Ambulatory Surgical Home On Campus Outpatient Hospital Office

Off Campus Outpatient Hospital
Pharmacy

Send completed form to: Case Review Unit CVS Caremark Specialty Programs Fax: 1-855-330-1720

Note: This fax may contain medical information that is privileged and confidential and is solely for the use of individuals named above. If you are not the intended recipient you hereby are advised that any dissemination, distribution, or copying of this communication is prohibited. If you have received the fax in error, please immediately notify the sender by telephone and destroy the original fax message. Myobloc MR SGM - 07/2022.

CVS Caremark Specialty Pharmacy • 2211 Sanders Road NBT-6 • Northbrook, IL 60062

Phone: 1-888-877-0518 • Fax: 1-855-330-1720 • www.caremark.com

Exception Criteria Questions:

- A. The preferred products for your patient's health plan are Botox, Dysport, and Xeomin. Can the patient's treatment be switched to one of the preferred products?
 □ Yes, *Please obtain Form for preferred product and submit for corresponding PA*. □ No
- B. Is this request for the treatment of cervical dystonia? \Box Yes \Box No If No, skip to D
- C. Has the patient had a documented inadequate response or intolerable adverse event to treatment with all of the preferred products (Botox, Dysport and Xeomin)? *Action Required: If 'Yes', Attach supporting chart note(s).*□ Yes □ No *Yes or No, skip to Clinical Criteria Questions*
- D. Is this request for the treatment of chronic sialorrhea? Yes No If No, skip to Criteria questions
- E. Has the patient experienced a documented inadequate response or intolerable adverse event to treatment with the preferred product (Xeomin)? *Action Required: If Yes, attach supporting chart note(s)* \Box Yes \Box No

<u>Clinical Criteria Questions:</u>

What is the diagnosis?
 Cervical dystonia (e.g., torticollis)
 Excessive salivation (chronic sialorrhea)
 Other

D Primary axillary or palmer hyperhidrosis

Upper limb spasticity

- 2. What is the ICD-10 code?
- 3. Is therapy prescribed for cosmetic purposes (e.g., treatment of wrinkles)? \Box Yes \Box No

Complete the following section based on the patient's diagnosis, if applicable.

Section A: Cervical Dystonia

- 4. Is the patient an adult? \Box Yes \Box No
- 5. Prior to initiating therapy with Myobloc, was/is there abnormal placement of the head with limited range of motion in the neck? Yes No

Section B: Excessive Salivation

6. Is the patient refractory to pharmacotherapy (for example, anticholinergics)? \Box Yes \Box No

Section C: Primacy Axillary or Palmer Hyperhidrosis

- Has significant disruption of professional and/or social life occurred because of excessive sweating?
 □ Yes □ No
- 8. Has the patient tried topical aluminum chloride or other extra-strength antiperspirants? \Box Yes \Box No
- 9. Was the topical aluminum chloride or other extra-strength antiperspirants ineffective or result in a severe rash?
 □ Yes □ No
- 10. Is the patient unresponsive or unable to tolerate oral pharmacotherapy prescribed for excessive sweating (e.g., anticholinergics, beta-blockers, or benzodiazepines)? □ Yes □ No

Section D: Upper Limb Spasticity

11. Is the spasticity a primary diagnosis or a symptom of a condition causing limb spasticity? \Box Yes \Box No

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Step Therapy Override: Complete if Applicable for the state of Maryland.		Please Circle	
Is the requested drug being used to treat stage four advanced metastatic cancer?	Yes	No	
Is the requested drug's use consistent with the FDA-approved indication or the National Comprehensive Cancer Network Drugs & Biologics Compendium indication for the treatment of stage four advanced metastatic cancer and is supported by peer-reviewed medical literature?	Yes	No	
Is the requested drug being used for an FDA-approved indication OR an indication supported in the compendia of current literature (examples: AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)?	Yes	No	
Does the prescribed quantity fall within the manufacturer's published dosing guidelines or within dosing guidelines found in the compendia of current literature (examples: package insert, AHFS, Lexicomp, Clinical Pharmacology, Micromedex, current accepted guidelines)?	Yes	No	
Do patient chart notes document the requested drug was ordered with a paid claim at the pharmacy, the pharmacy filled the prescription and delivered to the patient or other documentation that the requested drug was prescribed for the patient in the last 180 days?	Yes	No	
Has the prescriber provided proof documented in the patient chart notes that in their opinion the requested drug is effective for the patient's condition?	Yes	No	

Step Therapy Override: Complete if Applicable for the state of Virginia.		Please Circle	
Is the requested drug being used for an FDA-approved indication or an indication supported in the compendia of current literature (examples: AHFS, Micromedex, current accepted guidelines)?	Yes	No	
Does the prescribed dose and quantity fall within the FDA-approved labeling or within dosing guidelines found in the compendia of current literature?	Yes	No	
Is the request for a brand drug that has an AB-rated generic equivalent or interchangeable biological product available?	Yes	No	
Has the patient had a trial and failure of the AB-rated generic equivalent or interchangeable biological product due to an adverse event (examples: rash, nausea, vomiting, anaphylaxis) that is thought to be due to an inactive ingredient?	Yes	No	
Is the preferred drug contraindicated?	Yes	No	
Is the preferred drug expected to be ineffective based on the known clinical characteristics of the patient and the prescription drug regimen?	Yes	No	
Has the patient tried the preferred drug while on their current or previous health benefit plan and it was discontinued due to lack of efficacy or effectiveness, diminished effect, or an adverse event?		No	
Is the patient currently receiving a positive therapeutic outcome with the requested drug for their medical condition?	Yes	No	

I attest that this information is accurate and true, and that documentation supporting this information is available for review if requested by CVS Caremark or the benefit plan sponsor.

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Prescriber or Authorized Signature

Date (mm/dd/yy)

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