

BOTULINUM TOXINS PRIOR AUTHORIZATION FORM



Keystone First

PERFORMRxSM
Next Generation Pharmacy Benefits

(form effective 1/6/2025)

Fax to PerformRxSM at **1-866-497-1387**, or to speak to a representative call **1-800-588-6767**.

PRIOR AUTHORIZATION REQUEST INFORMATION

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request	Total # pages:	Name of office contact:
Contact's phone number:	LTC facility contact/phone:	

PATIENT INFORMATION

Patient name:	Patient ID #:	DOB:
Street address:	Apt #:	City/state/zip:

PRESCRIBER INFORMATION

Prescriber name:	Specialty:		
State license #:	NPI:	MA Provider ID #:	
Street address:	Suite #:	City/state/zip:	
Phone:	Fax:		

CLINICAL INFORMATION

Product requested: <input type="checkbox"/> Botox (preferred with clinical PA required) <input type="checkbox"/> Dysport (preferred with clinical PA required) <input type="checkbox"/> Myobloc (non-preferred) <input type="checkbox"/> Xeomin (non-preferred)		
Strength:	Injection site(s) and dose per site:	Qty requested:
Diagnosis (submit documentation):		DX code (required):

PHARMACY INFORMATION (Prescriber to identify the pharmacy that is to dispense the medication):

Deliver to: <input type="checkbox"/> Patient's Home <input type="checkbox"/> Physician's Office <input type="checkbox"/> Patient's Preferred Pharmacy Name:	
Pharmacy Phone #:	Pharmacy Fax #:
<input type="checkbox"/> I acknowledge that the patient agrees with the pharmacy chosen for delivery of this medication.	

INITIAL REQUESTS (Complete questions applicable to drug requested and patient's diagnosis):

- Request for a non-preferred agent (Myobloc or Xeomin):** Does the patient have a history of trial and failure, contraindication, or intolerance of the preferred Botulinum Toxins that are FDA-approved for the patient's diagnosis and age? Check all that apply. ☐ Botox ☐ Dysport
☐ Yes ☐ No ☐ N/A *Submit documentation of all medications tried and outcomes.*
- Axillary hyperhidrosis:** Does the patient have a history of trial and failure, contraindication, or intolerance of a topical agent such as 20% aluminum chloride?
☐ Yes ☐ No *List medications tried.*
- Overactive bladder:** Does the patient have a history of trial and failure, contraindication, or intolerance of at least two other medications used to treat OAB?
☐ Yes *List medication tried:* _____
☐ No
- Urinary incontinence due to detrusor overactivity associated with a neurologic condition:** Does the patient have a history of trial and failure, contraindication, or intolerance of at least one anticholinergic medication used to treat urinary incontinence? ☐ Yes ☐ No *List medications tried.*
- Migraine, Chronic:** Check all of the following that apply to the patient and submit documentation for each.

<input type="checkbox"/> Has a diagnosis of chronic migraine headache according to the current International Headache Society Classification of Headache Disorders that is not attributed to other causes including medication overuse	<input type="checkbox"/> History of trial and failure, contraindication, or intolerance of an agent in at least two of the following drug classes used for migraine prevention: <input type="checkbox"/> anticonvulsants <input type="checkbox"/> beta blockers <input type="checkbox"/> antidepressants
<input type="checkbox"/> The requested agent is prescribed by, or in consultation with, one of the following specialists. Submit documentation of consultation, if applicable. <input type="checkbox"/> neurologist <input type="checkbox"/> headache specialist who is certified in headache medicine by the United Council for Neurologic Subspecialties (UCNS)	<input type="checkbox"/> calcitonin gene-related peptide (CGRP)-targeting migraine preventive therapies <i>List medications tried:</i> _____
- Spasticity, Chronic:** Check all of the following that apply to the patient and submit documentation for each.

<input type="checkbox"/> has spasticity that interferes with activities of daily living <input type="checkbox"/> has spasticity that is expected to result in joint contracture with future growth
<input type="checkbox"/> if the patient has developed contractures, has been considered for surgical intervention
<input type="checkbox"/> if ≥ 18 years of age: <input type="checkbox"/> has focal spasticity <input type="checkbox"/> has tried and failed, or has contraindication or intolerance of, an oral medication for spasticity <i>List medications tried:</i> _____
<input type="checkbox"/> drug is being requested to either: <input type="checkbox"/> enhance function --OR-- <input type="checkbox"/> allow for additional therapeutic modalities to be employed
<input type="checkbox"/> drug will be used in conjunction with other appropriate therapeutic modalities (e.g., OT, PT, gradual splinting)
- All other diagnoses:** Submit documentation supporting the use of the requested agent for the patient's diagnosis and other treatments tried:

RENEWAL REQUESTS

Check all of the following that apply to the patient and submit documentation for each:

1. Request for frequency of injection that is consistent with the dose and duration of therapy limits:

☐ Patient showed a positive response to the medication

☐ For treatment of chronic migraine headache:

☐ Patient requires repeat injection to reduce the frequency, severity, or duration of symptoms

☐ The requested agent is prescribed by, or in consultation with, one of the following specialists. Submit documentation of consultation, if applicable.

☐ neurologist ☐ headache specialist who is certified in headache medicine by the United Council for Neurologic Subspecialties (UCNS)

☐ For treatment of all other diagnoses:

☐ Patient's symptoms returned to such a degree that repeat injection is required

2. Request for frequency of injection that exceeds the dose and duration of therapy limits:

☐ Treatment was well tolerated but inadequate.

☐ Peer-reviewed medical literature supports more frequent dosing as safe and effective for the diagnosis and requested dose (submit documentation of peer-reviewed medical literature)

PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION

Prescriber signature:	Date:
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