

**BOTULINUM TOXINS PRIOR AUTHORIZATION FORM** (form effective 1/6/2025)

Prior authorization guidelines for **Botulinum Toxins** and **Quantity Limits/Daily Dose Limits** are available on the DHS Pharmacy Services website at <https://www.pa.gov/en/agencies/dhs/resources/for-providers/ma-for-providers/pharmacy-services.html>.

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request		Total # of pages _____	Prescriber name:	
Name of office contact:			Specialty:	
Contact's phone number:			NPI:	State license #:
LTC facility contact/phone:			Street address:	
Beneficiary name:			City/state/zip:	
Beneficiary ID#:	DOB:	Phone:	Fax:	

**CLINICAL INFORMATION**

Drug requested:	Units/package size:	Total quantity requested per treatment:
Injection site(s) & dose per site:		
Diagnosis ( <u>submit documentation</u> ):	Dx code ( <u>required</u> ):	
Dates of previous administration and injection sites ( <u>submit documentation</u> ):		

**Complete all sections that apply to the beneficiary and this request.  
 Check all that apply and SUBMIT DOCUMENTATION for each item.**

**INITIAL requests**

- For a NON-PREFERRED Botulinum Toxin:**
  - Has a history of trial and failure of or a contraindication or an intolerance to the preferred Botulinum Toxins that are approved or medically accepted for treatment of the beneficiary's diagnosis (Refer to <https://papdl.com/preferred-drug-list> for a list of preferred and non-preferred drugs in this class.)
- For a diagnosis of CHRONIC SPASTICITY:**
  - Has spasticity that interferes with activities of daily living
  - Has spasticity that is expected to result in joint contracture with future growth
  - If the beneficiary has contractures, has been considered for surgical intervention
  - One of the following:
    - Has focal spasticity
    - Is under 18 years of age
    - Is 18 years of age or older and tried and failed or has a contraindication or an intolerance to an oral medication for spasticity

- Botulinum Toxin is prescribed to enhance function or allow for additional therapeutic modalities to be used
- Will use the requested botulinum toxin in conjunction with other appropriate therapeutic modalities (e.g., PT, OT, gradual splinting, etc.)

**For a diagnosis of AXILLARY HYPERHIDROSIS:**

- Tried and failed or has a contraindication or an intolerance to a topical agent such as aluminum chloride 20% solution

**For a diagnosis of CHRONIC MIGRAINE HEADACHE:**

- Has a diagnosis of migraine headache consistent with the current International Headache Society Classification of Headache Disorders
- Migraine headache is not attributable to other causes, such as medication overuse
- Is prescribed the Botulinum Toxin by or in consultation with a headache specialist who is certified in headache medicine by the United Council for Neurologic Subspecialties or a neurologist
- Tried and failed or has a contraindication or an intolerance to at least one drug used for migraine prevention from at least 2 of the following classes:
  - Anticonvulsants (e.g., divalproex, topiramate, valproic acid)
  - Antidepressants (e.g., amitriptyline, venlafaxine)
  - Beta blockers (e.g., metoprolol, propranolol, timolol)
  - CGRP-targeting migraine preventive therapies (e.g., gepants, monoclonal antibodies)

**For a diagnosis of URINARY INCONTINENCE due to detrusor overactivity:**

- Has an associated neurologic condition
- Tried and failed or has a contraindication or an intolerance to an anticholinergic drug used for the treatment of urinary incontinence (e.g., darifenacin, fesoterodine, oxybutynin, solifenacin, tolterodine, trospium)

**For a diagnosis of OVERACTIVE BLADDER:**

- Has symptoms of urge urinary incontinence, urgency, and frequency
- Tried and failed or has a contraindication or an intolerance to at least 2 drugs used for the treatment of overactive bladder (e.g., anticholinergics, beta-3 adrenergic agonists)

**RENEWAL requests**

- Experienced a positive clinical response to the Botulinum Toxin
- One of the following:
  - For the treatment of chronic migraine headache, requires repeat injection to reduce the frequency, severity, or duration of symptoms
  - For the treatment of all other diagnoses, has symptoms that returned to such a degree that repeat injection with Botulinum Toxin is required
- The frequency of injection of Botulinum Toxin exceeds the FDA-approved package labeling
  - The previous treatment was well-tolerated but inadequate
  - The requested dose and increased frequency of injection of Botulinum Toxin are supported by medical literature as safe and effective for the diagnosis
- For a diagnosis of CHRONIC MIGRAINE HEADACHE:**
  - Is prescribed the Botulinum Toxin by or in consultation with a headache specialist who is certified in headache medicine by the United Council for Neurologic Subspecialties or a neurologist

**PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION TO 866-240-3712**

Prescriber Signature:

Date:

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