Clinical Criteria Worksheet: OnabotulinumtoxinA (Botox®)

Enrollee Information	
Enrollee Last Name:	Enrollee First Name:
Date of Birth (MM/DD/YYYY):	Enrollee Medicaid ID (2 letters, 5 numbers, 1 letter):
City, Town or Post Office:	State: ZIP Code:
Prescriber Information	
Prescriber Last Name:	Prescriber First Name:
National Provider Identifier (NPI) Number: Preferred Contact (Telephone Number):	
Clinical Criteria – Drug Information Drug Administration: Provide the date of drug administration (MM/DD/YY / /	ΎΥ):
Drug name and strength:	
OnabotulinumtoxinA (Botox [®]) 200 units vial	
New treatment: Yes No	
If No , date therapy initiated (MM/DD/YYYY):	

Clinical Criteria – Diagnosis

1. Diagnosis related to use:

Food and Drug Administration Indications:	Compendia supported uses:
Primarly axillary hyperhidrosis (severe)	Achalasia
Blepharospasm associated with dystonia	Auricolotemporal syndrome
Cervical dystonia	Backache
Chronic migraine prophylaxis	Benign prostatic hyperplasia
Neurogenic detrusor overactivity	Cervicogenic headache
Overactive bladder	Difficulty talking – total laryngectomy
Spasticity	Dysphagia
Strabismus	Disorder of nervous system – excessive salivation
	Epicondylitis
	Excessive salivation – Advanced Parkinson's disease
	🗌 Fibromyalgia
	Gilles de la Tourette's syndrome
	Granuloma of vocal cords – refractory to conventional surgical/medical
	therapies
	Hemifacial spasm
	Idiopathic trigeminal neuralgia – refractory
	Injury to oculomotor nerve (acute)
	Isolated oromandibular dystonia
	Larynx closure – adjunct to surgical procedure
	Organic voice tremor
	Pelvic floor dyssynergia
	Spasm of pharyngoesophageal segment – total laryngectomy
	Spastic dysphonia
	Stuttering
	Tardive dyskinesia
	Temporomandibular joint disorder
	Whiplash injury to neck
Other:	

Clinical Criteria

2. Please indicate if this request is for the initiation or continuation of OnabotulinumtoxinA therapy:

Initiation Continuation

	month, with headache lasting greater than or equ	ual to 4 hours per day?	
	Yes No Not Applicable		
	d or Compendia supported oral preventive agents prolol, nadolol, propranolol, timolol], divalproex amate, or venlafaxine)?		
	Yes No Not Applicable		
	Please provide the names of the most recent therapies and dates of the trials. Please write N/A if thi question is not applicable based on diagnosis.		
	Drug name and strength:	Date(s) of use:	
	Drug name and strength:	Date(s) of use:	
4.	. If for overactive bladder or urinary incontinence due to detrusor overactivity, has the patient tried an antimuscarinic agent or beta-3 adrenergic agonist?		
	Yes No Not Applicable		
Please provide the name of the most recent therapy and dates of the trial. Please write N/ question is not applicable based on diagnosis.			

3. If for chronic migraine prophylaxis, does the patient have headaches greater than or equal to 15 days per

Drug name and strength: Date(s) of use:	
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5. If for neurogenic detrusor overactivity, does the patient have a diagnosis of multiple sclerosis or spinal cord injury?

Yes	No 🗌] Not Applicable
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If NO, has the patient tried an antimuscarinic agent?

Yes No Not Applicable

Please provide the name of the most recent therapy and dates of the trial. Please write N/A if this question is not applicable based on diagnosis.

Drug name and strength:______ Date(s) of use:______

Attestation

I attest that this drug is medically necessary for this patient and that all of the information on this form is accurate to the best of my knowledge. I attest that documentation of the above diagnosis and medical necessity is available for review if requested by the New York State Medicaid Program.

Prescriber Sign	ature (Required)
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Date (MM/DD/YYYY)