

Enrollee Last Name:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Enrollee First Name:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Clinical Criteria – Drug Information

Drug Administration:

Provide the date of drug administration (MM/DD/YYYY):

		/			/				
--	--	---	--	--	---	--	--	--	--

Drug name and strength:

- IncobotulinumtoxinA (Xeomin®) 50 units vial
- IncobotulinumtoxinA (Xeomin®) 100 units vial
- IncobotulinumtoxinA (Xeomin®) 200 units vial

Patient's current weight: _____ kg

Administration dose (units) and frequency: _____

Quantity of vials needed: _____

New treatment: Yes No

If No, date therapy initiated (MM/DD/YYYY):

		/			/				
--	--	---	--	--	---	--	--	--	--

Clinical Criteria – Diagnosis

1. Food and Drug Administration Indications:

- Blepharospasm
- Cervical dystonia
- Chronic sialorrhea
- Upper limb spasticity
- Other: _____

Enrollee Last Name:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Enrollee First Name:

--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--	--

Clinical Criteria

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

Initiation Continuation

3. If for chronic sialorrhea, has the patient had a trial with glycopyrrolate?

Yes No Not Applicable

If NO, does the patient have a diagnosis of Parkinson's disease or other neurodegenerative disease?

Yes No Not Applicable

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 Signature (Required)

Date (MM/DD/YYYY)