



Enrollee Last Name:

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Enrollee First Name:

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## Clinical Criteria – Drug Information

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### Drug Administration:

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

		/			/				
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### 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):

		/			/				
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## Clinical Criteria – Diagnosis

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### 1. Food and Drug Administration Indications:

- Blepharospasm  
 Cervical dystonia  
 Chronic sialorrhea  
 Upper limb spasticity  
 Other: \_\_\_\_\_

Enrollee Last Name:

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Enrollee First Name:

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### Clinical Criteria

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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

Archived December 2024

### Attestation

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*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.*

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Prescriber Signature (Required)

Date (MM/DD/YYYY)