

BOTOX (onabotulinumtoxinA)

Instructions

Please complete Part A and have your physician complete Part B. Completion and submission is not a guarantee of approval. Any fees related to the completion of this form are the responsibility of the plan member. Drugs in the Prior Authorization Program may be eligible for reimbursement if the patient does not qualify for coverage under a primary plan or a government program. Drugs used for indications not approved by Health Canada may be denied. For Quebec plan members, RAMQ exception drug criteria may apply. The decision for approval versus denial is based on pre-defined clinical criteria, primarily based on Health Canada approved indication(s) and on supporting evidence-based clinical protocols. The plan member will be notified whether their request has been approved or denied. Please note that you have the right to appeal the decision made by Express Scripts Canada.

Part A – Patient

First Name:		Last Name:		
Insurance Carrier Name/Number:				
Group Number:		Client ID:		
Date of Birth (YYYY/MM/DD):		Relationship: Employee Spouse Dependent		
Language: English French		Gender: Male Female		
Address:				
City:	Province:		Postal Code:	
Email address:				
Telephone (home):	Telephone (cell):		Telephone (work):	

Coordination of benefits

Patient Assistance	Is the patient enrolled in any patient assistance program?			
Program	Contact Name: Fax:			
Provincial Coverage	Has the patient applied for reimbursement under a provincial plan? Yes No N/A			
	What is the coverage decision of the drug? Approved Denied *Attach decision letter*			
Primary Coverage	Has the patient applied for reimbursement under a primary plan?			
	What is the coverage decision of the drug? Approved Denied *Attach decision letter*			

Authorization

On behalf of myself and my eligible dependents, I authorize my group benefit provider, and its agents, to exchange the personal information contained on this form. I give my consent on the understanding that the information will be used solely for purposes of administration and management of my group benefit plan. This consent shall continue so long as my dependents and I are covered by, or are claiming benefits under the present group contract, or any modification, renewal, or reinstatement thereof.

Plan Member Signature

Date



BOTOX (onabotulinumtoxinA)

Part B - Prescriber

Please see instructions on page 1 and complete all sections below. <u>Incomplete forms may result in automatic denial</u>. Please do **not** provide genetic test information or results.

SECTION 1 – DRUG REQUESTED

BOTOX (onabotulinu	imtoxinA)	New request	Renewal request*	
Dose	Administration (ex: oral, IV, e	tc) Frequency	Duration	
Site of drug administrat	ion:			
Home F	hysician's office/Infusion clinic	Hospital (outpatient)	Hospital (inpatient)	
* Please submit proof of prior coverage if available				

SECTION 2 – ELIGIBILITY CRITERIA

1. Please indicate if the patient satisfies the below criteria:
Blepharospasm
For the treatment of blepharospasm associated with dystonia, including benign essential blepharospasm, or VII nerve disorders (e.g. hemifacial spasm), AND
The patient is 12 years of age or older
Cervical Dystonia
For the treatment of cervical dystonia (spasmodic torticollis) in an adult
Focal Spasticity
For the treatment of focal spasticity, including upper limb spasticity in an adult, OR
For the treatment of lower limb spasticity in an adult, OR
For the treatment of upper and/or lower limb spasticity in pediatric patients 2 to 17 years of age
Strabismus
For the treatment of strabismus, AND
The patient is 12 years of age or older
Bladder Dysfunction – Neurogenic Detrusor Overactivity
For the treatment of urinary incontinence due to neurogenic detrusor overactivity resulting from neurogenic bladder associated multiple sclerosis or sub cervical spinal cord injury in an adult, AND
The patient has had an inadequate response or has a documented intolerance to anticholinergic medications (Please list prior therapies in the chart below)



BOTOX (onabotulinumtoxinA)

Bladder Dysfunction - Ov	veractive Bladder					
For the treatment of overactive bladder with symptoms of urinary incontinence, urgency, and frequency in an adult, AND						
The patient has had an inadequate response or has a documented intolerance to anticholinergic medications (<i>Please list prior therapies in the chart below</i>)						
Migraine						
For the prevention	on of migraine in an adult, AND					
The patient has	The patient has at least 15 headache days per month, AND					
	had an inadequate response or is es in the chart below), AND	intolerant to at least	1 prophylactic t	herapy for migrai	nes (Please	
BOTOX will not b	e used in combination with a calc	itonin gene-related pe	eptide (CGRP) aı	ntagonist		
Primary Hyperhidrosis of	the Axillae – Eligibility based on p	lan design				
For the treatmer	t of hyperhidrosis of the axillae in	an adult				
OR						
None of the above	ve criteria applies.					
Relevant additional information:						
	triad tharaniaa					
2. Please list previously						
Drug	Dosage and	Duration	Duration of therapy		Reason for cessation Inadequate Allergy/	
	administratio	n From	То	response	Intolerance	



BOTOX (onabotulinumtoxinA)

SECTION 3 - PRESCRIBER INFORMATION

Physician's Name:				
Address:				
Tel:		Fax:		
License No.:		Specialty:		
Physician Signature:		Date:		
Please fax or mail the completed form to Express Scripts Canada®	Fax: Express Scripts Canada Cl 1 (855) 712-6329	inical Services	Mail:	Express Scripts Canada Clinical Services 5770 Hurontario Street, 10 th Floor Mississauga, ON L5R 3G5