

Outpatient Medical Injectables Botulinum Toxin Request Form. Fax to 833-581-1861 (Medical Benefit Only)

Member Name:	DOB:	ID (UMI):	Medicare Commercial*	
Ordering/Attending Provider Name:		NPI Number	:	
Ordering/Attending Provider Address:				
Office Contact:	Phone #:	Fax #:_		
Servicing Facility/Vendor Name:	Facility NPI:			
Servicing Facility/Vendor Address:				
ICD10 Diagnosis Code(s):	Requested Start Date of Service:			
☐ Buy & Bill ☐ Drug Supplied by	Specialty Pharmacy (Pharmacy	Name:	NPI:	
ВОТОХ (Ј0585)			XEOMIN (J0588)	
OTHER	(J)			
Dose or number of units:	Frequency:	Number o	f visits requested:	
FOR CHRONIC MIGRAINE				
How many days a month does the mer	mber experience headache?			
When the member experiences migraines, how many hours a day do they last?				
For how long has the member been ex	periencing migraine headache	s?		
Is this request prescribed by or in cons	ultation with a neurologist or	headache specialist? YES	□ NO	
Is a healthcare provider trained in adm	inistration of botox administe	ring the drug? ☐ YES ☐ NO)	
Has the diagnosis of chronic migraine he Edition? (ICHD-III) ☐ YES ☐ NO				
Has there been a persistent three mon or calendar? ☐ YES ☐ NO	th history of recrurring debilit	ating headache documented b	y the member via headache diary	
Are headaches caused by medication r				
Has the member tried and failed adequate trials of prophylactic therapy from at least two different therapy classes (ex: antiseizure, beta blocker, tricyclic antidepressant)? \square YES \square NO				
Please list all previous prophylactic therapies tried and failed, not tolerated or contraindicated:				
Were the above medications prescribed at adequate doses for reasonable lengths of time (ex: 6 weeks each)? ☐ YES ☐ NO				

FOR CHRONIC MIGRAINE			
☐ New Start	☐ Continuation of Therapy		
	Since starting Botox has the member's migraine headache frequency reduced by at least 50% from baseline? ☐ YES ☐ NO Since starting Botox has the member's migraine headache hours reduced by at least 50% from baseline? ☐ YES ☐ NO		
Please attach all pertinent clinical information			
FOR HYPERHIDROSI			
Does the member have severe hyperidrosis? ☐ YES ☐ NO			
Please indicate which focal region the botulinum toxin will be treating: (circle all that apply)			
Axillary Region Palmar Region Plantar Region Craniofacial Region Other:			
Please indicate if the member has experienced any of the following:			
History of recurrent skin maceration with bacterial or fungal infections? □ YES □ NO			
 History of atopic dermatitis (atopic eczema) despite medical treatments with topical dermatological or systemic anticholinergic agents? ☐ YES ☐ NO 			
Has the member been unresponsive or unable to tolerate pharmacotherapy modalities prescribed for excessive sweating (example anticholinergics, beta-blockers, or benzodiazepines)? \square YES \square NO			
Have topical products such as 20% aluminum chloride or other extra strength antiperspirants been ineffective or resulted in a severe rash? \square YES \square NO			
☐ New Start	Continuation of Therapy		
	Since starting botulinum toxin, is there a documented objective measurable effect indicating a positive clinical response to treatment (ex: improvement in HDSS)?		
	☐ YES please describe: ☐ NO		
FOR ALL OTHER USES			

☐ New Start

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Has the member had a documented positive clinical response to treatment? ☐ YES ☐ NO

☐ YES ☐ NO

☐ Continuation of Therapy

All references to "Highmark" in this document are references to the Highmark company that is providing the member's health benefits or health benefit administration and/or to one or more of its affiliated Blue companies

Attached:

Please list all other therapies tried and failed, not tolerated, or contraindicated for the diagnosis:

^{**}Please verify member's eligibility and benefits through the health plan**