

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

Member Name:	DOB:	ID (UMI):	Medicare Commercial*
		NPI Number:	
Ordering/Attending Provider Address:_			
Office Contact:	Phone #:	Fa	x #:
Servicing Facility/Vendor Name:		Facility NPI:	
Servicing Facility/Vendor Address:			
ICD10 Diagnosis Code(s):		Requested Start Date of Service:	
*For providers in Western PA and West Virgin	ia, the specialty pharmacy will be a	ssigned by Free Market Health	. All other providers, please specify below:
☐ Buy & Bill ☐ Drug Supplied by	Specialty Pharmacy (Pharmacy	v Name:	NPI:
ВОТОХ (Ј0585)			XEOMIN (J0588)
OTHER	(J)		
Dose or number of units:	Frequency:	Numb	er of visits requested:
FOR CHRONIC MIGRAINE			
How many days a month does the member experience headache?			
When the member experiences migraines, how many hours a day do they last?			
For how long has the member been experiencing migraine headaches?			
Is this request prescribed by or in consultation with a neurologist or headache specialist? NO			
Is a healthcare provider trained in adm		ring the drug? ☐ YES ☐	NO
Has the diagnosis of chronic migraine Edition? (ICHD-III) ☐ YES ☐ NO			
Has there been a persistent three mor or calendar? ☐ YES ☐ NO	nth history of recrurring debilit	ating headache document	ed by the member via headache diary
Are headaches caused by medication r	-		
		erapy from at least two di	fferent therapy classes (ex: antiseizure,
 beta blocker, tricyclic antidepressant)? ☐ YES ☐ 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			

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

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		
FOR HYPERHIDROSIS			
Does the member have severe hyperidrosis?			
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 (ex: 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? YES 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			
Please list all other therapies tried and failed, not tolerated, or contraindicated for the diagnosis:			
☐ New Start	☐ New Start ☐ Continuation of Therapy		
	Has the member had a documented positive clinical response to treatment? ☐ YES ☐ NO		
Please attach all pertinent clinical information			
	Attached: YES NO		

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