SENTARA COMMUNITY PLAN (MEDICAID)

PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST*

<u>Directions</u>: <u>The prescribing physician must sign and clearly print name (preprinted stamps not valid)</u> on this request. All other information may be filled in by office staff; <u>fax to 1-800-750-9692</u>. No additional phone calls will be necessary if all information (<u>including phone and fax #s</u>) on this form is correct. <u>If the information provided is not complete, correct, or legible, the authorization process can be delayed.</u>

Botulinum Toxin Injections®, Type A (Pharmacy)

<u>Drug Requested</u>: **Botox**[®] (onabotulinumtoxinA)

(Chronic Migraine Headache Prophylaxis)

MEMBER & PRESCRIBER INFORMATION	Authorization may be delayed if incomplete.
Member Name:	
Member Sentara #:	Date of Birth:
Prescriber Name:	
Prescriber Signature:	
Office Contact Name:	
Phone Number:	Fax Number:
NPI #:	
DRUG INFORMATION: Authorization may be de	elayed if incomplete.
Drug Name/Form/Strength:	
Dosing Schedule:	
Diagnosis:	ICD Code, if applicable:
Weight (if applicable):	Date weight obtained:

- Max quantity limits: 155 units once every 12 weeks
- Cosmetic indications are <u>EXCLUDED</u>

NOTE: In treating adult patients for one or more indications, the maximum cumulative dose should not exceed 400 units, in a 3-month interval. In pediatric patients, the total dose should not exceed the lower of 10 units/kg body weight or 340 units, in a 3-month interval.

CLINICAL CRITERIA: Check below all that apply. <u>All criteria must be met for approval</u>. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

Diagnosis: Chronic Migraine Headache Prophylaxis		
Initial Authorization Approval: 12 months		
	Has the member been approved for Botox previously through the Sentara Health Plan medical department?	
	\square Member must be ≥ 18 years of age	
	☐ Member experiences ≥ 15 headache days per month	
	\square Member experiences headaches which last ≥ 4 hours per day	
	Member must have failed a <u>2-month</u> trial of at least one medication from <u>TWO (2)</u> different migraine prophylactic classes supported by American Headache Society/American Academy of Neurology treatment guidelines 2012/2015/2021, Level A and B evidence; ICSI 2013, high quality evidence (verified by pharmacy paid claims or submitted chart notes):	
	☐ Anticonvulsants (divalproex, valproate, topiramate)	
	☐ Beta blockers (atenolol, metoprolol, nadolol, propranolol, timolol)	
	☐ Antidepressants (amitriptyline, venlafaxine)	
	 □ Injectable CGRP inhibitors (Aimovig[®], Emgality[®], Ajovy[®]) or oral CGRP inhibitors indicated for migraine prevention (Qulipta[™], Nurtec ODT[™]) * requires prior authorization* 	
	Member has been evaluated for medication overuse headache (MOH) (defined as headaches occurring greater than or equal to 15 days per month. It develops as a consequence of regular overuse of acute of symptomatic headache medication for more than 3 months)	
	Treatment will include a plan to taper off the offending medication if MOH is diagnosed	
appr	uthorization Approval: 12 months. Check below all that apply. All criteria must be met for oval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart, must be provided or request may be denied.	
	Member has experienced a positive response to therapy, demonstrated by a reduction in headache frequency	
	Use of acute migraine medications (e.g., NSAIDs, triptans) has decreased since the start of Botox®	
	Member continues to be monitored for medication overuse headache (MOH)	
Medication being provided by (check box below that applies):		
	Physician's office OR Specialty Pharmacy – PropriumRx	

**Use of samples to initiate therapy does not meet step edit/preauthorization criteria. **

*Previous therapies will be verified through pharmacy paid claims or submitted chart notes. *