# 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 (including phone and fax  $\#_s$ ) on this form is correct. <u>If the information provided is not</u> complete, correct, or legible, the authorization process can be delayed.

## **Botulinum Toxin Injections<sup>®</sup>**, Type A (Pharmacy)

**Drug Requested: Botox**<sup>®</sup> (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: Authoriza | ation may be delayed if incomplete. |
| Drug Name/Form/Strength:    |                                     |
| Dosing Schedule:            | Length of Therapy:                  |
| 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.

(Continued on next page)

#### **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? □ Yes
  - $\Box \quad \text{Member must be} \ge 18 \text{ years of age}$
  - $\Box$  Member experiences  $\geq 15$  headache days per month
  - $\Box$  Member experiences headaches which last  $\geq 4$  hours per day
- □ Member must have failed a 2-month trial of at least one medication from TWO (2) 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<sup>®</sup>, Emgality<sup>®</sup>, Ajovy<sup>®</sup>) or oral CGRP inhibitors indicated for migraine prevention (Qulipta<sup>TM</sup>, Nurtec ODT<sup>TM</sup>) \* 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 or symptomatic headache medication for more than 3 months)
- □ Treatment will include a plan to taper off the offending medication if MOH is diagnosed

Reauthorization Approval: 12 months. Check below all that apply. <u>All criteria must be met for</u> approval. To support each line checked, all documentation, including lab results, diagnostics, and/or chart notes, must be provided or request may be denied.

- □ Member has experienced a positive response to therapy, demonstrated by a reduction in headache frequency
- $\Box$  Use of acute migraine medications (e.g., NSAIDs, triptans) has decreased since the start of Botox<sup>®</sup>
- □ 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.\*

 $\Box$  No