Botox Injection for Chronic Migraine Protocol

**Purpose/Indication:** Prevention of headaches in adults (18 years or older) with chronic migraine who have 15 or more headaches each month.

**Definition of chronic migraine:** Per the International Headache Society (IHS), chronic migraine is defined as a condition in which headaches occur on 15 or more days per month for more than 3 months, which, on at least 8 days per month, has features of migraine headache. The term “episodic migraine” is used to describe migraine headaches with a frequency of 14 or fewer days per month.

**Population Foci (CRNP):** Adult Acute Care, Adult Health, Adult/Gerontology Acute Care, Adult/Gerontology Primary Care, Family, and Gerontology.

**CRNP or PA:** ___________________________ License # __________

**Physician Attestation (s):**

- Supervising/collaborating physician trained in Botox injections for chronic migraines with at least 1 year experience performing Botox injections for chronic migraine pursuant to FDA approved protocols.

- The collaborating/supervising or covering physician will be physically available on site.

- Prior to approval for performing injections, the NP/PA will receive three hours of didactic training, attend a half day course with a physician preceptor certified by the pharmaceutical company.

- NP/PA must observe 10 procedures and perform 10 procedures under direct supervision. Following approval, the NP/PA must perform no less than 25 treatments per year to maintain competency. (1 set of injections equals 1 procedure)

- The collaborating/supervising or covering physician will be responsible for diagnosing and referring the patient for Botox treatment. NP/PA may only perform treatment on patients with a physician diagnosis of chronic migraine (i.e., headache occurs 15 or more days per month for three months).

- Prior to each treatment, the NP/PA will evaluate for contraindications by visually inspecting the injection sites, palpating the injection sites, and asking the patient to activate the muscle at the injection sites. The treatment is approximately 15 minutes and involves a 0.1 ml injection into 31 standardized sites on the patient’s head and neck. The dosage and injection sites are the same for all patients receiving the treatment.

- Prior to providing the treatment, the NP/PA must discuss with the collaborating/supervising physician whether the patient is receiving one of the enumerated drugs that potentially interferes with neuromuscular transmission or whether the patient has pre-existing neuromuscular disorders. In addition, the treatment by the NP/PA is contraindicated if the injection site is infected or if the patient has a hypersensitivity to Botox.

- On a quarterly basis, four patients are to be randomly chosen for discussion with the collaborating/supervising physician. The NP/PA and physician will immediately discuss any adverse events.
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Description of skill/ procedure in detail:

1. Patient condition/ exam finding
   a. Patient must have a diagnosis of Chronic Migraine, as defined on page 1, by a physician
2. FDA Approved Treatment Schedule
   a. Fixed dosing of Onabotulinumtoxin A every 12 weeks
3. Pre-examination assessment of brow, forehead, and neck to evaluate the patient for conditions that may be affected or exacerbated by treatment. If any conditions are present, the NP/PA should inform the physician prior to continuing treatment.
   a. Visually inspect the muscle
   b. Ask the patient to activate the muscle
   c. Palpate the muscle
4. Onabotulinumtoxin A injection sites and order of injections using 0.1 mL (5 units) of Onabotulinumtoxin A per site. The procedure takes approximately 15 minutes to administer 31 injections in 7 head and neck muscle areas.
   a. Corrugator
      i. 10 units divided in 2 sites
   b. Procerus
      i. 5 units in 1 site
   c. Frontalis
      i. 20 units divided in 4 sites
   d. Temporalis
      i. 40 units divided in 8 sites
   e. Occipitalis
      i. 30 units divided in 6 sites
   f. Cervical paraspinals
      i. 20 units divided in 4 sites
   g. Trapezius
      i. 30 units divided in 6 sites
5. Expected results
   a. A decrease in the frequency from baseline of headache days
6. After care
   a. Report any adverse reactions to provider
7. Follow up
   a. Fixed dose schedule every 12 weeks

No deviation from FDA approved protocol including dosage, location, and number of injections is allowed.

Physician Signature: ___________________________________________ License # _______________________

Physician Name (Print): ___________________________________________ Date: ______________

Page 2 of 2

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