

# SENTARA HEALTH PLANS

## PHARMACY PRIOR AUTHORIZATION/STEP-EDIT REQUEST\*

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

### Injectable Calcitonin Gene-Related Peptide (CGRP) Antagonists

**Drug Requested:** (Select one from below)

PREFERRED	
<input type="checkbox"/> <b>Aimovig<sup>®</sup></b> (erenumab)	<input type="checkbox"/> <b>Emgality<sup>®</sup></b> (galcanezumab)
NON-PREFERRED	
<input type="checkbox"/> <b>Ajovy<sup>®</sup></b> (fremanezumab) *Member must have tried and failed <b><u>BOTH</u></b> preferred agents and meet all PA criteria for approval of Ajovy*	

**Sentara Considers the use of concomitant therapy with Calcitonin Gene-Related Peptide Antagonists (CGRP) and Botox to be experimental and investigational, although safety and efficacy of these combinations has been established. In the event a member has an active Botox authorization on file and dual therapy is requested, all subsequent CGRP requests will be reviewed and assessed for medical necessity of combination therapy.**

**MEMBER & PRESCRIBER INFORMATION:** Authorization may be delayed if incomplete.

Member Name: \_\_\_\_\_

Member Sentara #: \_\_\_\_\_ Date of Birth: \_\_\_\_\_

Prescriber Name: \_\_\_\_\_

Prescriber Signature: \_\_\_\_\_ Date: \_\_\_\_\_

Office Contact Name: \_\_\_\_\_

Phone Number: \_\_\_\_\_ Fax Number: \_\_\_\_\_

DEA OR NPI #: \_\_\_\_\_

**DRUG INFORMATION:** Authorization may be delayed if incomplete.

Drug Form/Strength: \_\_\_\_\_

Dosing Schedule: \_\_\_\_\_ Length of Therapy: \_\_\_\_\_

Diagnosis: \_\_\_\_\_ ICD Code: \_\_\_\_\_

Weight: \_\_\_\_\_ Date: \_\_\_\_\_

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**Recommended Dosing & Quantity Limits:**

Drug	Dose	Quantity Limit
<b>Aimovig®</b> (erenumab)	<ul style="list-style-type: none"> <li><b>Migraine Prophylaxis:</b> Initial: 70 mg SC once a month; some members may benefit from 140 mg once a month (given as 2 consecutive 70 mg injections)</li> </ul>	<ul style="list-style-type: none"> <li>70 mg/mL (1 mL/30 day)</li> <li>140 mg dose (2 mL/30 days)</li> <li>If using the 140 mg dose, must use the package labeled specifically for 140 mg/mL</li> </ul>
<b>Ajovy®</b> (fremanezumab)	<ul style="list-style-type: none"> <li><b>Migraine Prophylaxis:</b> 225 mg SC monthly <b>or</b> 675 mg every 3 months</li> </ul>	<ul style="list-style-type: none"> <li>225 mg/1.5 mL; 1.5 mL (1 syringe) per 30 days or 4.5 mL (3 syringes) per 90 days</li> </ul>
<b>Emgality®</b> (galcanezumab)	<ul style="list-style-type: none"> <li><b>Migraine Prophylaxis:</b> Initial: 240 mg SC as a single loading dose, followed by 120 mg once monthly</li> <li><b>Episodic cluster headache prophylaxis:</b> 300 mg SC at the onset of the cluster period and then once monthly until the end of the cluster period</li> </ul>	<ul style="list-style-type: none"> <li>120 mg/mL; 1 mL (1 auto-injector and prefilled syringe) per 30 days with one time loading dose of 2 mL (2 auto-injectors)</li> <li>For Episodic Cluster headache diagnosis only: 300 mg dose; 100 mg/mL prefilled syringe</li> </ul>

**CLINICAL CRITERIA:** Check below all that apply. All criteria must be met for approval.

**Authorization Criteria**

- ☐ Member must be 18 years of age or older
- ☐ Provider has attested to all clinical criteria for **ONE** of the applicable diagnoses below

**DIAGNOSIS:** Please check **ONE** of the applicable diagnoses below

☐ **Chronic & Episodic Migraine Headache Prevention** (All applicable boxes below must be met to qualify)

- ☐ Member must have a diagnosis of Chronic or Episodic Migraine Headache defined by **BOTH** of the following:
  - ☐ Member has  $\geq 4$  migraine headache days per month
  - ☐ Member must have failed a **2-month** trial of at least one medication from **TWO** different migraine prophylactic classes supported by the American Headache Society/American Academy of Neurology treatment guidelines 2012/2015/2021, Level A and B evidence; ICSI 2013, high quality evidence:
    - ☐ Anticonvulsants (divalproex, valproate, topiramate)
    - ☐ Beta blockers (atenolol, metoprolol, nadolol, propranolol, timolol)
    - ☐ Antidepressants (amitriptyline, venlafaxine)

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- ☐ Member will **NOT** be initiating botulinum toxin headache prophylaxis after starting the requested agent
- ☐ Requested medication will **NOT** be used in combination with Botox or another CGRP inhibitor indicated for migraine prevention
- ☐ **For Ajovy Requests:** Member must have tried and failed **BOTH** preferred agents Aimovig and Emgality **AND** meet all prior authorization criteria for approval of Ajovy

☐ **Episodic Cluster Headaches (Emgality® Only)** (All applicable boxes below must be met to qualify)

- ☐ Member has between one headache every other day and eight headaches per day
- ☐ Member must have failed at least a **1-month** trial of at least **ONE** **generic** standard prophylactic pharmacologic therapy, used to prevent cluster headache and supported by the American Headache Society/American Academy of Neurology treatment guidelines:
  - ☐ Suboccipital steroid injection
  - ☐ Calcium channel blockers (verapamil)
  - ☐ Alkali metal/ Antimanic (lithium)
  - ☐ Anticoagulant (warfarin)
  - ☐ Anticonvulsants (topiramate)

***\*\*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.\****