

# OPTIMA HEALTH COMMUNITY CARE AND OPTIMA FAMILY CARE (MEDICAID)

## 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 will be delayed.**

### ANTIMIGRAINE DRUGS-OTHERS

#### [Calcitonin Gene-Related Peptide (CGRP) Antagonists]

**Drug Requested – check below drug that applies:**

#### PREFERRED Drugs (Require a prior authorization)

**Emgality<sup>®</sup>** (galcanezumab-gnlm) **Syringe**       **Emgality<sup>®</sup>** (galcanezumab-gnlm) **Pen**

#### Non-Preferred Drugs

**Aimovig<sup>®</sup>** (erenumab-aooe) Injection       **Ajovy<sup>®</sup>** (fremanezumab-vfrm) Injection

**DRUG INFORMATION:** Complete **all** information below or authorization will be delayed.

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

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

**Diagnosis:** \_\_\_\_\_ **ICD Code, if applicable:** \_\_\_\_\_

**Recommended dosage** (for subcutaneous use only):

**Emgality<sup>®</sup> Dosage:** 240 mg loading dose (administered as 2 consecutive injections of 120 mg each), followed by monthly doses of 120 mg.

**Aimovig<sup>®</sup> Dosage:** 70 mg once a month; some patients may benefit from 140 mg once a month (given as 2 consecutive 70 mg injections). **Quantity Limit:** 1 autoinjector/30 days; 2 autoinjectors/30days; 2 prefilled syringes/ 30 days

**Ajovy<sup>®</sup> Dosage:** 225 mg monthly or 675 mg quarterly dosage administered as 3 consecutive injections of 225 mg each; single-dose prefilled syringe – 225 mg/1.5mL solution

**CLINICAL CRITERIA:** Check below **ALL** that apply. **ALL** criteria **must** be met for approval. **ALL** documentation, including lab results and/or chart notes (**when required**), **must** be provided or request will be denied.

**Initial Approval – three (3) months**

**Does the member meet the following criteria?**

1. Does the member have a diagnosis of migraine with or without aura based on International Classification of Headache Disorders (ICHD-III) diagnostic criteria?  Yes     No

**AND**

(Continued on next page)

2. Is member 18 years or older?  Yes  No

**AND**

3. Member does not have medication over-use headache (MOH)?  Yes  No

**AND**

4. Women of childbearing age have had a pregnancy test at baseline?  Yes  No

**AND**

5. Member has  $\geq 4$  migraine days per month for at least 3 months?  Yes  No

**AND**

6. Member is utilizing prophylactic intervention modalities (e.g., **behavioral therapy, physical therapy, or life-style modifications**)?  Yes  No

**AND**

7. Member has tried and failed a  $\geq 1$  month trial of any **TWO (2)** of the **following** oral medications?  Yes  No

- Antidepressants (e.g., **amitriptyline, venlafaxine**)
- Beta blockers (e.g., **propranolol, metoprolol, timolol, atenolol**)
- Anti-epileptics (e.g., **valproate, topiramate**)
- Angiotensin converting inhibitors/angiotensin II receptor blockers (e.g., **lisinopril, candesartan**)

**Renewal Approval – Twelve (12) months:** **ALL** boxes below **must** be checked to ensure authorization will be **NOT** delayed.

8. Did member demonstrate significant decrease in the number, frequency, and/or intensity of headaches?  Yes  No

**AND**

9. Does member have an overall improvement in function with therapy?  Yes  No

**AND**

10. Does member continue to utilize prophylactic intervention modalities (e.g., **behavioral therapy, physical therapy, life-style modification**)?  Yes  No

**AND**

11. Women of childbearing age continue to be monitored for pregnancy status?  Yes  No

**AND**

12. Absence of unacceptable toxicity (e.g., **intolerable injection site pain or constipation**)?  Yes  No

(Continued on next page; signature **MUST** be attached with this request.)

**\*\*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.\***

Patient Name: \_\_\_\_\_

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

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

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

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

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

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

\*Approved by Pharmacy and Therapeutics Committee: 7/19/2018

REVISED/UPDATED: 9/30/2018; 11/3/2018; 2/17/2019; (Reformatted) 4/11/2019; 6/19/2019