AvMed

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 1-305-671-0200</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>

Injectable Calcitonin Gene-Related Peptide (CGRP) Antagonists

Drug Requested: (Select one from below)

PREFERRED				
	Aimovig [®] (erenumab)	□ Emgality [®] (galcanezumab)		
NON-PREFERRED				
	Ajovy [®] (fremanezumab) *Member must have tried and failed <u>BOTH</u> preferred agents and meet all PA criteria for approval of Ajovy*			

The Health Plan 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 AvMed #:	Date of Birth:	
Prescriber Name:		
Prescriber Signature:		
Office Contact Name:		
Phone Number:		
DEA OR NPI #:		
DRUG INFORMATION: Authorization may be delayed	l if incomplete.	
Drug Form/Strength:		
Dosing Schedule:	Length of Therapy:	
Diagnosis:	ICD Code:	
Weight: Date:	:	

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Drug	Dose	Quantity Limit
Aimovig [®] (erenumab)	• Migraine Prophylaxis : Initial: 70 mg SC once a month; some members may benefit from 140 mg once a month (given as 2 consecutive 70 mg injections)	 70 mg/mL (1 mL/30 day) 140 mg dose (2 mL/30 days) If using the 140 mg dose, must use the package labeled specifically for 140 mg/mL
Ajovy [®] (fremanezumab)	• Migraine Prophylaxis: 225 mg SC monthly or 675 mg every 3 months	• 225 mg/1.5 mL; 1.5 mL (1 syringe) per 30 days or 4.5 mL (3 syringes) per 90 days
Emgality [®] (galcanezumab)	 Migraine Prophylaxis: Initial: 240 mg SC as a single loading dose, followed by 120 mg once monthly Episodic cluster headache prophylaxis: 300 mg SC at the onset of the cluster period and then once monthly until the end of the cluster period 	 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) For Episodic Cluster headache diagnosis only: 300 mg dose; 100 mg/mL prefilled syringe

Recommended Dosing & Quantity Limits:

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

<u>Authorization Criteria</u>

- □ Member must be 18 years of age or older
- □ Provider has attested to all clinical criteria for <u>ONE</u> of the applicable diagnoses below

DIAGNOSIS: Please check <u>ONE</u> 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 **<u>BOTH</u>** of the following:
 - $\Box \quad \text{Member has} \ge 4 \text{ migraine headache days per month}$
 - □ Member must have failed a <u>2-month</u> trial of at least one medication from <u>TWO</u> different migraine prophylactic classes supported by the American Headache Society/American Academy of Neurology treatment guidelines 2012/2015/2021/2024, 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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- □ Angiotensin II receptor blocker (candesartan) *requires prior authorization*
- □ Injectable CGRP inhibitors (Aimovig[®], Emgality[®], Ajovy[®]) or oral CGRP inhibitors indicated for migraine prevention (Qulipta[™], Nurtec ODT[®]) *requires prior authorization*
- □ Member will <u>NOT</u> be initiating botulinum toxin headache prophylaxis after starting the requested agent
- □ Requested medication will <u>NOT</u> be used in combination with Botox or another CGRP inhibitor indicated for migraine prevention
- □ For Ajovy[®] Requests: Member must have tried and failed <u>BOTH</u> preferred agents Aimovig[®] and Emgality[®] <u>AND</u> 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 <u>1-month</u> trial of at least <u>ONE</u> 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. *<u>Previous therapies will be verified through pharmacy paid claims or submitted chart notes.</u>*