

CGRP Antagonists and Gepants Drug Use Criteria

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Table 1: FDA-Approved Indications for CGRP Antagonists

Drug	FDA-Approved Indication
Atogepant (Qulipta®)	Preventative migraine treatment
Eptinezumab (Vyapti®)	Preventative migraine treatment
Erenumab (Aimovig®)	Preventative migraine treatment
Fremanezumab (Ajovy®)	Preventative migraine treatment
Galcanezumab (Emgality®)	Preventative migraine treatment and cluster headache prevention
Rimegepant sulfate (Nurtec ODT®)	Acute migraine treatment and preventative treatment of episodic migraine
Ubrogepant (Ubrelvy®)	Acute migraine treatment
Zavegepant (Zavzpret®)	Acute migraine treatment

Preferred agents (if criteria met for approval):

- ubrogepant (Ubrelvy®) – for acute (abortive) therapy
- fremanezumab (Ajovy®) and erenumab (Aimovig®) – for preventative therapy

GUIDELINE FOR USE:

Initial Authorization

1. Does the member have a diagnosis of migraine?
 - a. If yes, go to 2
 - b. If no, deny as criteria not met. Off-label use of a medication is not a covered benefit under OHP

2. Do chart notes indicate headaches are due to medication overuse (MOH)?
 - a. If yes, send for MD review to determine medical appropriateness.
 - b. If no, got to 3

3. Is the request for acute (abortive) migraine treatment AND the patient is 18 years or older?
 - a. If yes, go to 9
 - b. If no, go to 4

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4. Is the request for prevention of cluster headache and the patient is 18 years or older?
 - a. If yes, go to 11
 - b. If no, go to 5

 5. Is the request for prophylactic therapy and the patient is 18 years or older?
 - a. If yes, go to 6
 - b. If no, deny as criteria not met

 6. Is there is documentation that the patient has experienced 4 or more migraine days in the previous month
 - a. If yes, go to 7
 - b. If no, deny as criteria not met

 7. Has the patient had an adequate trial (2-6 months) without response, or has contraindications, to at least THREE of the following OHP preferred drugs (in the same or different classes)?
 - Propranolol immediate-release, metoprolol, or atenolol
 - Topiramate, valproic acid, or divalproex sodium
 - Amitriptyline, nortriptyline, or venlafaxine
 - Candesartan or telmisartan
 - OR does the patient have a documented intolerance, FDA-labeled contraindication or hypersensitivity to the above migraine prophylaxis medications?
 - a. If yes, go to 8
 - b. If no, deny as criteria not met

 8. Has the patient received an injection with botulinum toxin for headache treatment once in the previous 2 months?
 - a. If yes, deny as criteria not met. Concurrent use of a CGRP antagonist and Botox is not a covered benefit.
 - b. If no, approve for up to 6 months

 9. In a patient with acute migraines, has the patient failed to receive benefit from adequate trial with at least TWO different triptans or have contraindications to triptans?
 - a. If yes, go to 10
 - b. If no, deny as criteria not met. Previous trial of at least two preferred formulary triptans is required.

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10. Does the patient have a history of at least 4 migraines a month AND is on preventative migraine therapy (excluding other CGRP inhibitors)?
- If yes, approve for up to 6 months
 - If no, deny as criteria not met
11. Has the patient had an adequate trial (2-6 months) without response, or has contraindications, to at least THREE of the following:
- Lithium
 - Verapamil
 - Suboccipital steroid injection
 - Sumatriptan subcutaneous
 - Zolmitriptan nasal spray
- If yes, approve for 6 months
 - If no, deny as criteria not met

Renewal Authorization

- Do chart notes indicate headaches are due to medication overuse?
 - If yes, send for MD review to determine medical appropriateness
 - If no, go to 2
- Has the patient experienced a documented positive response to therapy, as demonstrated by a reduction in migraine/cluster headache frequency and/or intensity from baseline?
 - If yes, approve for 12 months
 - If no, deny as criteria not met

Rationale: To ensure step therapy and appropriate use of high-cost migraine therapies aligned with clinical guidelines and OHP expectations.

References:

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