



Prior Authorization
<p>JOHNS HOPKINS HEALTHCARE (MEDICAID) Self-administered CGRP Inhibitors - Priority Partners MCO</p> <p>This fax machine is located in a secure location as required by HIPAA regulations. Complete/review information, sign and date. Fax signed forms to Johns Hopkins Healthcare at 1-410-424-4607. Please contact Johns Hopkins Healthcare at 1-888-819-1043 with questions regarding the Prior Authorization process. When conditions are met, we will authorize the coverage of Self-administered CGRP Inhibitors - Priority Partners MCO.</p>

Drug Name (select from list of drugs shown)		
Aimovig (erenumab-aooe)	Ajovy (fremanezumab-vfrm)	Emgality (galcanezumab-gnlm)

Quantity	Frequency	Strength
Route of Administration	Expected Length of Therapy	

Patient Information	
Patient Name:	_____
Patient ID:	_____
Patient Group No.:	_____
Patient DOB:	_____
Patient Phone:	_____

Prescribing Physician	
Physician Name:	_____
Physician Phone:	_____
Physician Fax:	_____
Physician Address:	_____
City, State, Zip:	_____

Diagnosis: _____	ICD Code: _____
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Comments: _____

Please circle the appropriate answer for each question.	
1. Is this request for continuation of therapy?	<input type="checkbox"/> Y <input type="checkbox"/> N
[Note: The use of physician samples, or manufacturer product discounts, does not guarantee coverage under the provisions of the medical and/or pharmacy benefit. All pertinent criteria must be met in order to be eligible for benefit coverage.]	
[If no, then skip to question 4.]	
2. Has the patient's mean monthly headache days been reduced by 50 percent or greater relative to the patient's baseline prior to treatment?	<input type="checkbox"/> Y <input type="checkbox"/> N

[Note: Documentation must be submitted.]	
[If yes, then no further questions.]	
3. Has the patient had a reduction in any of the following validated migraine-specific patient-reported outcome measures: A) Migraine Disability Assessment (MIDAS): Reduction of 5 points or greater when baseline score is 11-20 OR reduction of 30 percent or greater when baseline score is greater than 20, B) Migraine Physical Function Impact Diary (MPFID): Reduction of 5 points or greater, C) Headache Impact Test (HIT-6): Reduction of 5 points or greater?	<input type="checkbox"/> Y <input type="checkbox"/> N
[Note: Documentation must be submitted.]	
[No further questions.]	
4. Is this request for concurrent use with medical botulinum toxin injection OR concurrent use with another calcitonin gene-related peptide (CGRP) agent?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If yes, then no further questions.]	
5. Is the patient 18 years of age or older?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If no, then no further questions.]	
6. Is the requested drug being prescribed by or in consultation with a neurologist or pain specialist?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If no, then no further questions.]	
7. Is this request for preventative chronic migraine management?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If no, then skip to question 10.]	
8. Does the patient have a documented diagnosis of chronic migraine (greater than or equal to 15 headache days per month with 8 being migraine days with symptoms lasting 4 hours a day or longer)?	<input type="checkbox"/> Y <input type="checkbox"/> N
[Note: Documentation must be submitted.]	
[If no, then no further questions.]	
9. Has the prescriber documented that the patient does not have medication-induced headaches from overuse of acute treatment agents (analgesics, triptans, ergots)?	<input type="checkbox"/> Y <input type="checkbox"/> N
[Note: Documentation must be submitted.]	
[If yes, then skip to question 12.]	
[If no, then no further questions.]	
10. Is this request for episodic migraine management?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If no, then skip to question 15.]	
11. Does the patient have a documented diagnosis of episodic migraine (less than 15 days per month)?	<input type="checkbox"/> Y <input type="checkbox"/> N
[Note: Documentation must be submitted.]	
[If no, then no further questions.]	

12. Has the patient tried and failed medications from at least TWO of the following therapeutic classes used in standard therapy for migraine prophylaxis: A) Beta antagonists, B) Anticonvulsants, C) Antidepressants?	<input type="checkbox"/> Y <input type="checkbox"/> N
[Note: Documentation must be submitted.]	
[If no, then no further questions.]	
13. Is this request for Ajovy?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If yes, then no further questions.]	
14. Has the patient had a documented trial and failure with Ajovy?	<input type="checkbox"/> Y <input type="checkbox"/> N
[Note: Documentation must be submitted.]	
[No further questions.]	
15. Is this request for Emgality 100mg/ml?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If no, then no further questions.]	
16. Is this request for the treatment of episodic cluster headache?	<input type="checkbox"/> Y <input type="checkbox"/> N
[If no, then no further questions.]	
17. Does the patient have a documented diagnosis of episodic cluster headache (cluster headache attacks in periods lasting from 7 days to one year, separated by pain-free periods lasting at least 3 months)?	<input type="checkbox"/> Y <input type="checkbox"/> N
[Note: Documentation must be submitted.]	
[If no, then no further questions.]	
18. Has the prescriber documented that the patient has a maximum of 8 attacks per day, and a minimum of one attack every other day?	<input type="checkbox"/> Y <input type="checkbox"/> N
[Note: Documentation must be submitted.]	
[If no, then no further questions.]	
19. Has the patient tried and failed verapamil and at least one of the following: A) Prednisone, B) Dexamethasone?	<input type="checkbox"/> Y <input type="checkbox"/> N
[Note: Documentation must be submitted.]	

I attest that the medication requested is medically necessary for this patient. I further attest that the information provided is accurate and true, and that the documentation supporting this information is available for review if requested by the claims processor, the health plan sponsor, or, if applicable a state or federal regulatory agency.

Prescriber (Or Authorized) Signature and Date