

MIGRAINE PREVENTION AGENTS PRIOR AUTHORIZATION FORM

(form effective 1/6/2025)



Fax to PerformRxSM at **1-888-981-5202**, or to speak to a representative call **1-866-610-2774**.

PRIOR AUTHORIZATION REQUEST INFORMATION

<input type="checkbox"/> New request <input type="checkbox"/> Renewal request	# of pages:	Name of office contact:
Contact's phone number:	LTC facility contact/phone:	

PATIENT INFORMATION

Patient name:	Patient ID #:	DOB:
Street address:	Apt. #:	City/state/zip:

PRESCRIBER INFORMATION

Prescriber name:	Specialty:	
State license #:	NPI:	MA Provider ID#:
Street address:	Suite #:	City/state/zip:
Phone:	Fax:	

PHARMACY INFORMATION (Prescriber to identify the pharmacy that is to dispense the medication):

Deliver to: <input type="checkbox"/> Patient's Home <input type="checkbox"/> Physician's Office <input type="checkbox"/> Patient's Preferred Pharmacy Name:	
Pharmacy Phone #:	Pharmacy Fax #:
<input type="checkbox"/> I acknowledge that the patient agrees with the pharmacy chosen for delivery of this medication.	

CLINICAL INFORMATION

Product requested (clinical prior auth required):		
Preferred	Non-Preferred	
<input type="checkbox"/> Aimovig 70 mg/ml autoinjector <input type="checkbox"/> Aimovig 140 mg/ml autoinjector <input type="checkbox"/> Ajovy 225 mg/1.5 ml autoinjector <input type="checkbox"/> Ajovy 225 mg/1.5 ml syringe	<input type="checkbox"/> Emgality 120 mg/ml autoinjector <input type="checkbox"/> Emgality 120 mg/ml syringe <input type="checkbox"/> Emgality 300 mg (100 mg/ml syringe x 3) <input type="checkbox"/> Nurtec ODT 75 mg <input type="checkbox"/> Qulipta Tablet 10 mg <input type="checkbox"/> Qulipta Tablet 30 mg <input type="checkbox"/> Qulipta Tablet 60 mg <input type="checkbox"/> Vyepti IV Solution 100 mg/ml <input type="checkbox"/> Other: _____	
Dose/directions	Quantity:	Refills:
Diagnosis (submit documentation):	DX code (required):	
Is the medication being prescribed by, or in consultation with, a neurologist or a headache specialist who is certified in headache medicine by the United Council for Neurologic Subspecialties (UCNS)? <input type="checkbox"/> Yes <i>Submit documentation of consultation, if applicable.</i> <input type="checkbox"/> No		

ALL INITIAL REQUESTS

- If the patient is currently using a Migraine Prevention Agent, one of the following:
☐ Will discontinue use of that Migraine Prevention Agent prior to starting the requested Migraine Prevention Agent
☐ Has a medical reason for concomitant use of both Migraine Prevention Agents that is supported by peer-reviewed literature or national treatment guidelines. Please explain:
- For a non-preferred agent: Does the patient have history of therapeutic failure, contraindication, or intolerance to the preferred CGRP monoclonal antibodies (mAbs) approved or medically accepted for their indication?
☐ Yes ☐ No
If yes, select medications tried.
☐ Aimovig ☐ Ajovy ☐ Emgality ☐ Nurtec ODT ☐ Other:

INITIAL REQUESTS FOR MIGRAINES

- Has the patient averaged 4 or more migraine days per month over the past 3 months? ☐ Yes ☐ No
- For gepant (e.g., Nurtec ODT, Qulipta): Does the patient have history of therapeutic failure, contraindication, or intolerance to the preferred CGRP monoclonal antibodies (mAbs) approved or medically accepted for their indication?
☐ Yes ☐ No
If yes, select medications tried.
☐ Aimovig ☐ Ajovy ☐ Emgality ☐ Other:
- Does the patient have a confirmed diagnosis of migraine (with or without aura) according to the current International Headache Society Classification of Headache Disorders?
☐ Yes ☐ No
- Does the patient have a history of trial and failure of or contraindication or intolerance to at least one drug from one of the following three classes?
☐ anticonvulsants (e.g., divalproex, topiramate, valproic acid) ☐ antidepressants (e.g., amitriptyline, venlafaxine) ☐ beta blockers (e.g., metoprolol, propranolol, timolol)
☐ Yes - List medications tried:
☐ No
- Provide average number of migraine days and headache days per month at baseline:



INITIAL REQUESTS FOR EPISODIC CLUSTER HEADACHE

1. Does the patient have confirmed diagnosis of episodic cluster headache according to the current International Headache Society Classification of Headache Disorders?
☐ Yes ☐ No
2. Does the patient have a history of trial and failure, contraindication, or intolerance of a preventive medication recommended by current consensus guidelines for episodic cluster headaches?
☐ Yes - *List medications tried:*
☐ No

RENEWAL REQUESTS

1. For the prevention of migraine: Since starting the requested medication, did the patient experience one of the following:
☐ Reduction in the average number of migraine days per month from baseline
☐ Decrease in severity or duration of migraines from baseline
2. For episodic cluster headache: Since starting the requested medication, did the patient experience a reduction in cluster headache frequency from baseline? ☐ Yes ☐ No
3. For gepant (e.g., Nurtec ODT, Qulipta), for the prevention of migraine: Does the patient have history of therapeutic failure, contraindication, or intolerance to the preferred CGRP monoclonal antibodies (mAbs) approved or medically accepted for their indication? ☐ Yes ☐ No
 If yes, select medications tried:
☐ Aimovig ☐ Ajovy ☐ Emgality ☐ Other:
4. For a non-preferred agent: Does the patient have history of therapeutic failure, contraindication, or intolerance to the preferred CGRP monoclonal antibodies (mAbs) approved or medically accepted for their indication? ☐ Yes ☐ No
 If yes, select medications tried:
☐ Aimovig ☐ Ajovy ☐ Emgality ☐ Nurtec ODT ☐ Other:

PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION

Prescriber signature:	Date:
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