## **MIGRAINE PREVENTION AGENTS** PRIOR AUTHORIZATION FORM





(form effective 1/6/2025)

Fax to PerformRx<sup>SM</sup> at **1-855-851-4058**, or to speak to a representative call **1-866-907-7088**.

PRIOR AUTHORIZATION REQ	UEST INFORMATION							
☐ New request ☐ Renewal request	# of pages:	Name of o	ame of office contact:					
Contact's phone number:	LTC facility	acility contact/phone:						
PATIENT INFORMATION								
Patient name:			Patient ID #:			DOB:		
Street address:		Apt.	#:	City/state/zij	):			
PRESCRIBER INFORMATION								
Prescriber name:			Specialty:		1			
State license #: NPI:					MA Provider ID#:	MA Provider ID#:		
Street address:			te #: City/state/zip:					
Phone:			Fax:					
<b>PHARMACY INFORMATION</b> (Prescriber to identify the pharmacy that is to dispense the medication):								
Deliver to:  Patient's Home Physician's Office Patient's Preferred Pharmacy Name:								
Pharmacy Phone #:				Pharmacy Fax #:				
☐ 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					
☐ Aimovig 70 mg/ml autoinjector☐ Aimovig 140 mg/ml autoinjector☐	<ul> <li>☐ Emgality 120 mg/ml autoinje</li> <li>☐ Emgality 120 mg/ml syringe</li> </ul>		☐ Qulipta Tablet 10 mg ☐ Qulipta Tablet 30 mg					
☐ Ajovy 225 mg/1.5 ml autoinjector	☐ Emgality 300 mg (100 mg/ml		☐ Qulipta Tablet 60 mg					
☐ Ajovy 225 mg/1.5 ml syringe	☐ Nurtec ODT 75 mg		U vyepti iv s	Solution 100 m	g/mi 			
Dose/directions			I		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)? ☐ Yes Submit documentation of consultation, if applicable. ☐ No								
ALL INITIAL REQUESTS								
1. If the patient is currently using a Migraine Prevention Agent, one of the following:								
☐ 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:								
2. For a non-professed agent: Dece the nations have history of the appealing failure, contraindication, or intelegence to the professed CCDD managinal antihodics (make) approved or								
2. 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								
1. Has the patient averaged 4 or more migraine days per month over the past 3 months? ☐ Yes ☐ No								
2. 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.								
if yes, select filedications thed.  □ Aimovig □ Ajovy □ Emgality □ Other:								
3. Does the patient have a confirmed diagnosis of migraine (with or without aura) according to the current International Headache Society Classification of Headache Disorders?								
4. 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								
5. Provide average number of migraine days and headache days per month at baseline:								

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INITIAL REQUESTS FOR EPISODIC CLUSTER HEADACHE						
. Does the patient have confirmed diagnosis of episodic cluster headache according to the current International Headache Society Classification of Headache Disorders?						
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?  \[ \subseteq \text{Yes} - \text{List medications tried:} \] \[ \subseteq \text{No} \]						
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   Since starting the requested medication, did the patient experience a reduction in cluster headache frequency from baseline?   Yes   No   No   No   No   No   No   No   No						
PLEASE FAX COMPLETED FORM WITH REQUIRED CLINICAL DOCUMENTATION						
Prescriber signature:	Date:					

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