Print Clear

Prior Authorization Request Administrative Information

Member Information

Last name _____ First name _____ MI _____

Member ID _____ Date of birth _____

Sex assigned at birth ☐ Female ☐ Male ☐ "X" or Intersex

Current gender ☐ Female ☐ Male ☐ Transgender male ☐ Transgender female ☐ Other _____

Place of residence ☐ Home ☐ Nursing facility ☐ Other _____

Race/ethnicity _____ Preferred spoken language _____ Preferred written language _____

MassHealth does not exclude people or treat them differently because of race, color, national origin, age, disability, religion, creed, sexual orientation, or sex (including gender identity and gender stereotyping).

Plan Contact Information

Please indicate the member's MassHealth Plan and fax or submit this completed and signed form according to the Plan's contact information below.

MassHealth Fee-For-Service (FFS) Plan, Primary Care Clinician (PCC) Plan, Primary Care Accountable Care Organization (PCACO) Plan, Children's Medical Security Plan, and Health Safety Net Plan
☐ **MassHealth Drug Utilization Review Program** Pharmacy: Fax: (877) 208-7428 - Tel: (800) 745-7318
MassHealth Managed Care Organization (MCO) and Accountable Care Partnership Plans (ACPP)
☐ **Fallon Health** Online Prior Authorization: go.covermymeds.com/OptumRx Online Prior Authorization: providerportal.surescripts.net/ProviderPortal/optum Pharmacy: Fax: (844) 403-1029 - Tel: (844) 720-0033
☐ **Health New England** Online Prior Authorization: go.covermymeds.com/OptumRx Pharmacy: Fax: (800) 550-9246 - Tel: (800) 918-7545
☐ **Mass General Brigham Health Plan** Online Prior Authorization (Non-Specialty Drugs): go.covermymeds.com/OptumRx Online Prior Authorization (Specialty/Medical Drugs): provider.massgeneralbrighamhealthplan.org Pharmacy: Fax: (844) 403-1029 - Tel: (800) 711-4555
☐ **Tufts Health Plan** Online Prior Authorization: point32health.promptpa.com Pharmacy: Fax: (617) 673-0939 - Tel: (888) 257-1985
☐ **WellSense Health Plan** Online Prior Authorization: wellsense.org/providers/ma/pharmacy/prior-authorizations Pharmacy: Fax: (833) 951-1680 - Tel: (877) 417-1822

over

Headache Therapy (Calcitonin Gene-Related Peptide [CGRP] Inhibitors) Prior Authorization Request

MassHealth reviews requests for prior authorization (PA) on the basis of medical necessity only. If MassHealth approves the request, payment is still subject to all general conditions of MassHealth, including current member eligibility, other insurance, and program restrictions. MassHealth will notify the requesting provider and member of its decision. Keep a copy of this form for your records. If faxing this form, please use black ink.

Additional information about these agents, including PA requirements and preferred products, can be found within the MassHealth Drug List at **www.mass.gov/druglist.**

Medication information

Medication requested

- ☐ Aimovig (erenumab-aooe)
- ☐ Ajovy (fremanezumab-vfrm)
- ☐ Emgality (galcanezumab-gnlm)
- ☐ Nurtec (rimegepant)
- ☐ Qulipta (atogepant)
- ☐ Ubrelvy (ubrogepant)
- ☐ Vyepti (eptinezumab-jjmr)[MB]
- ☐ Zavzpret (zavegepant)

Dose, frequency, and duration of medication requested

[MB] *This drug is available through the health care professional who administers the drug or in an outpatient or inpatient hospital setting. MassHealth does not pay for this drug to be dispensed through the retail pharmacy. If listed, prior authorization does not apply through the hospital outpatient and inpatient settings. Please refer to 130 CMR 433.408 for prior authorization requirements for other health care professionals. Notwithstanding the above, this drug may be an exception to the unified pharmacy policy; please refer to respective MassHealth Accountable Care Partnership Plans (ACPPs) and Managed Care Organizations (MCOs) for prior authorization status and criteria, if applicable.*

Indication (Check all that apply or include ICD-10 code, if applicable.)

- ☐ Cluster headache
- ☐ Migraine headache
  - ☐ Prophylaxis. Frequency of migraine attacks (days/month)
  - ☐ Acute treatment. Frequency of migraine attacks (number/month)
- ☐ Other

Please indicate billing preference. ☐ Pharmacy ☐ Prescriber in-office ☐ Hospital outpatient

If applicable, please also complete section for professionally administered medications at end of form.

Drug NDC (if known) or service code

Section I. Please complete for Aimovig, Ajovy, Emgality, Nurtec, Qulipta, and Vyepti requests for migraine prophylaxis.

1. For all requests except Nurtec, has the member had a trial with a beta-blocker (atenolol, metoprolol, nadolol, propranolol, timolol)?
   ☐ Yes. Please list the drug name, dates/duration of use, and outcomes below.*

   Drug name Dates/duration of use

   Did the member experience any of the following? ☐ Adverse reaction ☐ Inadequate response ☐ Other

Briefly describe details of adverse reaction, inadequate response, or other.

☐ No. Please explain why not.

2. For Aimovig, Ajovy, and Emgality requests, please document a trial of topiramate, a tricyclic antidepressant, valproic acid, or venlafaxine. For Qulipta and Vyepti requests, please document a trial of Botox, topiramate, a tricyclic antidepressant, valproic acid, or venlafaxine. Alternatively, provide clinical rationale for use of Aimovig, Ajovy, Emgality, Qulipta, or Vyepti instead of these agents.
   ☐ Yes. Please list the drug names, dates/duration of use, and outcomes below.*

   Drug name Dates/duration of use

   Did the member experience any of the following? ☐ Adverse reaction ☐ Inadequate response ☐ Other

   Briefly describe details of adverse reaction, inadequate response, or other.

   ☐ No. Please explain why not.
3. For Aimovig requests, has the member had a trial with Ajovy or Emgality?
   ☐ Yes. Please list the dates/duration of use and outcomes below. *

   Dates/duration of use

   Did the member experience any of the following? ☐ Adverse reaction ☐ Inadequate response ☐ Other

   Briefly describe details of adverse reaction, inadequate response, or other.

   ☐ No. Please explain why not.
4. For Nurtec, Qulipta, and Vyepti requests, please document a trial of Aimovig, Ajovy, or Emgality. Alternatively, provide clinical rationale for use of Nurtec, Qulipta, or Vyepti instead of these agents.
   ☐ Yes. Please list the drug names, dates/duration of use, and outcomes below.*

   Drug name Dates/duration of use

   Did the member experience any of the following? ☐ Adverse reaction ☐ Inadequate response ☐ Other

   Briefly describe details of adverse reaction, inadequate response, or other.

   ☐ No. Please explain why not.

Section II. Please complete for Nurtec and Ubrelvy requests for acute treatment of migraine.

1. Has the member had a trial with two triptans?
   ☐ Yes. Please list the drug names, dates/duration of use, and outcomes below.*

   Drug name Dates/duration of use

   Did the member experience any of the following? ☐ Adverse reaction ☐ Inadequate response ☐ Other

   Briefly describe details of adverse reaction, inadequate response, or other.

   Drug name Dates/duration of use

   Did the member experience any of the following? ☐ Adverse reaction ☐ Inadequate response ☐ Other

   Briefly describe details of adverse reaction, inadequate response, or other.

   ☐ No. Please explain why not.

over

For requests for quantities above 16 units/30 days for Nurtec and Ubrelvy, is the member currently receiving prophylaxis?

☐ Yes. Please specify.

Drug name Dose and frequency

Drug name Dose and frequency

☐ No. Please explain why prophylaxis is not appropriate for this member.

**Attach a letter with additional information regarding medication trials as applicable.*

Section III. Please complete for recertification requests for Emgality for cluster headache.

1. Is the member still actively having a cluster headache? ☐ Yes. ☐ No.
2. Has the member been initiated on prophylactic therapy for the cluster headache?

   ☐ Yes. Please specify.

   Drug name Dose and frequency

   Drug name Dose and frequency

   ☐ No. Please explain why prophylaxis is not appropriate for this member.

Section IV. Please complete for Zavzpret requests for acute treatment of migraine.

1. Has the member had a trial with two triptan nasal sprays?

   ☐ Yes. Please list the drug names, dates/duration of use, and outcomes below.*

   Drug name Dose and frequency

   Drug name Dose and frequency

   ☐ No. Please explain why triptan nasal sprays are not appropriate for this member.

2. Please describe medical necessity for the use of the requested dosage formulation.

Section V. Please complete and provide documentation for exceptions to Step Therapy.

1. Is the alternative drug required under the step therapy protocol contraindicated, or will likely cause an adverse reaction in, or physical or mental harm to the member? ☐ Yes ☐ No

   If yes, briefly describe details of contraindication, adverse reaction, or harm.

2. Is the alternative drug required under the step therapy protocol expected to be ineffective based on the known clinical characteristics of the member and the known characteristics of the alternative drug regimen?

   ☐ Yes ☐ No

   If yes, briefly describe details of known clinical characteristics of member and alternative drug regimen.

over

3. Has the member previously tried the alternative drug required under the step therapy protocol, or another alternative drug in the same pharmacologic class or with the same mechanism of action, and such alternative drug was discontinued due to lack of efficacy or effectiveness, diminished effect, or an adverse event?
   ☐ Yes ☐ No
   If yes, please provide details for the previous trial.
   Drug name _____ Dates/duration of use _____
   Did the member experience any of the following? ☐ Adverse reaction ☐ Inadequate response
   Briefly describe details of adverse reaction or inadequate response.
   _____
   _____
4. Is the member stable on the requested prescription drug prescribed by the health care provider, and switching drugs will likely cause an adverse reaction in or physical or mental harm to the member?
   ☐ Yes. Please provide details. _____
   ☐ No

Please continue to next page and complete Prescriber and Provider Information section.

over

Prior Authorization Request Prescriber and Provider Information

Prescriber Information

Last name* First name* MI

NPI* Individual MH Provider ID

DEA No. Office Contact Name

Address City State Zip

Email address

Telephone No.* Fax No.*

** Required*

Please also complete for professionally administered medications, if applicable.

Start date End date

Servicing prescriber/facility name ☐ Same as prescribing provider

Servicing provider/facility address

Servicing provider NPI/tax ID No.

Name of billing provider

Billing provider NPI No.

Is this a request for recertification? ☐ Yes ☐ No

CPT code No. of visits J code No. of units

Prescribing provider's attestation, signature, and date

I certify under the pains and penalties of perjury that I am the prescribing provider identified in the Prescriber information section of this form. Any attached statement on my letterhead has been reviewed and signed by me. I certify that the medical necessity information (per 130 CMR 450.204) on this form is true, accurate, and complete, to the best of my knowledge. I understand that I may be subject to civil penalties or criminal prosecution for any falsification, omission, or concealment of any material fact contained herein.

Prescribing provider's signature _____

Printed name of prescribing provider Date

(The form can either be signed by hand and then scanned, or it can be signed electronically using DocuSign or Adobe Sign. For electronic signatures, the signer can upload a picture of their wet signature. The typed text of a signature is not an acceptable form of an electronic signature.)