



Service Authorization (SA) Form

Antimigraine Agents, Others

If the following information is not complete, correct, or legible, the SA process can be delayed.

Please use one form per member.

MEMBER INFORMATION

Last Name: _____

First Name: _____

Medicaid ID Number: _____

Date of Birth: _____

Weight in Kilograms: _____

PRESCRIBER INFORMATION

Last Name: _____

First Name: _____

NPI Number: _____

Phone Number: _____

Fax Number: _____

DRUG INFORMATION

Drug Name/Form: _____

Strength: _____

Dosing Frequency: _____

Length of Therapy: _____

Quantity per Day: _____

Preventive treatment of migraine	
Preferred Agents *step edit required	Non-Preferred Agents (SA required)
Aimovig®, Ajovy® and Ajovy® autoinjector Emgality® pen and syringe (120 mg), Nurtec® ODT, Qulipta™	Emgality® syringe (100 mg), Vyepti®
Acute treatment of migraine	
Preferred Agents (No SA with trial of 2 generic triptans)	Non-Preferred Agents (SA required)
Nurtec® ODT, Ubrelvy™	Reyvow®, Trudhesa™, Zavzpret™

(Form continued on next page.)

Member's Last Name:

Member's First Name:

DRUG INFORMATION (*Continued*)

Identify why the preferred agents cannot be used.

DIAGNOSIS AND MEDICAL INFORMATION

All drugs in this class are eligible to receive a SIX (6)-month approval. Complete the following questions.

For Preventive treatment of migraine, does the member meet the *step edit AND the following criteria?

1. Does the member have a diagnosis of migraine with or without aura based on International Classification of Headache Disorders (ICHD-III) diagnostic criteria? **AND**
☐ Yes ☐ No
2. Is the member ≥ 18 years of age? **AND**
☐ Yes ☐ No
3. Has the member had ≥ 4 migraine days per month for at least 3 months? **AND**
☐ Yes ☐ No
4. *Has the member tried and failed a ≥ 1 month trial of any 2 of the following oral generic medications?
 - Antidepressants (e.g., amitriptyline, venlafaxine)
 - Beta blockers (e.g., propranolol, metoprolol, timolol, atenolol)
 - Anti-epileptics (e.g., valproate, topiramate)
 - Angiotensin converting enzyme inhibitors/angiotensin II receptor blockers (e.g., lisinopril, candesartan)☐ Yes ☐ No
5. For Nurtec and Qulipta, has the member tried and failed 1 of the preferred injectable agents?
☐ Yes ☐ No

For renewal, complete the following question to receive a TWELVE (12)-month approval.

1. Did the member demonstrate significant decrease in the number, frequency, or intensity of headaches?
☐ Yes ☐ No

(Form continued on next page.)

Member's Last Name:

Member's First Name:

For Acute treatment of migraine, does the member meet the *step edit AND the following criteria?

1. Does the member have a diagnosis of migraine with or without aura? **AND**

☐ Yes ☐ No

2. Is the member ≥ 18 years of age? **AND**

☐ Yes ☐ No

3. *Has the member tried and failed (or has contraindications to) two preferred triptan medications?

☐ Yes ☐ No

4. Prior to initiation of Trudhesa™, a cardiovascular evaluation is recommended. Has this been completed?

☐ Yes ☐ No

For renewal, complete the following question to receive a TWELVE (12)-month approval.

2. Did the member demonstrate significant decrease in the number, frequency, or intensity of headaches?

☐ Yes ☐ No

(Form continued on next page.)

Member's Last Name:

Member's First Name:

For Episodic Cluster Headache, does the member meet the following criteria?

1. Does the member have a diagnosis of episodic cluster headache? **AND**

☐ Yes ☐ No

2. Is the member ≥ 18 years of age? **AND**

☐ Yes ☐ No

3. Has the member experienced at least 2 cluster periods lasting from 7 days to 365 days, separated by pain-free periods lasting at least three months? **AND**

☐ Yes ☐ No

4. Medication will not be used in combination with another CGRP antagonist or inhibitor used for the preventive treatment of migraines? **AND**

☐ Yes ☐ No

5. Has the member tried and failed (or has contraindications to) at least one standard prophylactic (preventive) pharmacologic therapy for cluster headache?

☐ Yes ☐ No

For renewal, complete the following question to receive a TWELVE (12)-month approval.

1. Did the member demonstrate significant decrease in the number, frequency, or intensity of headaches?

☐ Yes ☐ No

Prescriber Signature (Required)

Date

By signature, the physician confirms the above information is accurate and verifiable by member records.

Please include ALL requested information; Incomplete forms will delay the SA process.

Submission of documentation does NOT guarantee coverage by the Department of Medical Assistance Services.

The completed form may be: **FAXED TO 800-932-6651**, phoned to 800-932-6648, or mailed to:

Prime Therapeutics Management LLC/Attn: GV – 4201

P.O. Box 64811, St. Paul, MN 55164-0811