



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

(Form continued on next page.)

Member's Last Name:

Member's First Name:

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**DRUG INFORMATION (Continued)**

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Identify why the preferred agents cannot be used.

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**DIAGNOSIS AND MEDICAL INFORMATION**

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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  $\geq 18$  years of age? **AND**

Yes     No

3. Has the member had  $\geq 4$  migraine days per month for at least 3 months? **AND**

Yes     No

4. \*Has the member tried and failed a  $\geq 1$  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:**

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**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  $\geq 18$  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:

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**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  $\geq$  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

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**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.

Fax this form to 1-866-940-7328

Pharmacy PA call center: 1-800-310-6826