

**FORWARDHEALTH
PRIOR AUTHORIZATION DRUG ATTACHMENT FOR HEADACHE AGENTS,
PREVENTATIVE TREATMENT**

INSTRUCTIONS: Type or print clearly. Before completing this form, read the Prior Authorization Drug Attachment for Headache Agents, Preventative Treatment Instructions, F-02667A. Prescribers may refer to the Forms page of the ForwardHealth Portal at www.forwardhealth.wi.gov/WIPortal/Subsystem/Publications/ForwardHealthCommunications.aspx?panel=Forms for the completion instructions.

Pharmacy providers are required to have a completed Prior Authorization Drug Attachment for Headache Agents, Preventative Treatment form signed and dated by the prescriber before submitting a prior authorization (PA) request on the Portal, by fax, or by mail. Prescribers and pharmacy providers may call Provider Services at 800-947-9627 with questions.

SECTION I – MEMBER INFORMATION

1. Name – Member (Last, First, Middle Initial)

2. Member ID Number

3. Date of Birth – Member

SECTION II – PRESCRIPTION INFORMATION

4. Drug Name

5. Drug Strength

6. Date Prescription Written

7. Refills

8. Directions for Use

9. Name – Prescriber

10. Address – Prescriber (Street, City, State, Zip+4 Code)

11. Phone Number – Prescriber

12. National Provider Identifier – Prescriber

SECTION III – CLINICAL INFORMATION – ALL REQUESTS

13. Diagnosis Code and Description

Note: Supporting clinical information and a copy of the member's current medical records must be submitted with all PA requests for non-preferred headache agents, preventative treatment drugs. Medical records must demonstrate that the member meets the clinical criteria and document the member's medical work-up for migraines, including the current number of headache days per month, the number of migraine days per month, the average migraine duration (in hours), as well as complete problem and medication lists.



SECTION IV – CLINICAL INFORMATION – INITIAL REQUESTS ONLY

14. Has the prescriber evaluated and diagnosed the member as having a history of migraines, with or without aura, according to the International Classification of Headache Disorders, third edition, diagnostic criteria? Yes No

15. Document the member's current headache frequency.

Headache Days Per Month _____ Migraine Days Per Month _____

Average Migraine Duration in Hours _____

16. Document the member's current migraine prescribed medication treatment regimen.

List the current prescribed migraine preventative medications (drug name[s], dose, and dosing frequency), **including Botox (if applicable)**.

List the current prescribed migraine rescue medications (drug name[s], dose, and dosing frequency).

17. Indicate the preferred headache agents, preventative treatment drugs the member has taken and provide specific details regarding the member's response to treatment and the reason(s) for discontinuing. If additional space is needed, continue documentation in Section VII of this form.

1. Drug Name _____ Dose _____ Dates Taken _____

Description of Treatment Response and Reason(s) for Discontinuing

2. Drug Name _____ Dose _____ Dates Taken _____

Description of Treatment Response and Reason(s) for Discontinuing

3. Drug Name _____ Dose _____ Dates Taken _____

Description of Treatment Response and Reason(s) for Discontinuing

SECTION V – CLINICAL INFORMATION – RENEWAL REQUESTS ONLY

18. Has the member experienced/sustained a clinically significant decrease in the number of migraine days per month and/or a decrease in migraine duration compared to their baseline prior to initiation of treatment with a headache agent, preventative treatment drug? Yes No

19. Document the member's current headache frequency.

Headache Days Per Month _____ Migraine Days Per Month _____

Average Migraine Duration in Hours _____

20. List the current prescribed migraine preventative medications (drug name[s], dose, and dosing frequency), **including Botox (if applicable)**.

List the current prescribed migraine rescue medications (drug name[s], dose, and dosing frequency).

Has the member been compliant with the current prescribed migraine medication treatment regimen? Yes No

SECTION VI – AUTHORIZED SIGNATURE

21. **SIGNATURE** – Prescriber

22. Date Signed

SECTION VII – ADDITIONAL INFORMATION

23. Include any additional information in the space below. Additional diagnostic and clinical information explaining the need for the drug requested may be included here.
