

Migraine Acute Treatment Agents
Phone: 215-991-4300
Fax back to: 866-240-3712

Health Partners Plans manages the pharmacy drug benefit for your patient. Certain requests for coverage require review with the prescribing physician. Please answer the following questions and fax this form to the number listed above.

PLEASE NOTE: Any information (patient, prescriber, drug, labs) left blank, illegible, or not attached WILL DELAY the review process.

Member Name:	Prescriber Name:	
HPP Member Number:	Fax:	Phone:
Date of Birth:	Office Contact:	
Member Primary Phone:	NPI:	PA PROMISE ID:
Address:	Address:	
City, State ZIP:	City, State ZIP:	
Line of Business: <input type="checkbox"/> Medicaid <input type="checkbox"/> CHIP	Specialty Pharmacy (if applicable):	
Drug Name:	Strength:	
Quantity:	Refills:	
Directions:		
Diagnosis Code:	Diagnosis:	
<i>HPP's maximum approval time is 12 months but may be less depending on the drug.</i>		

Please attach any pertinent medical history including labs and information for this member that may support approval.

Please answer the following questions and sign.

Q1. The request is for a Migraine Acute Treatment Agent that was previously approved. If YES, go to 17.

Yes No

Q2. The request is for a gepant for the preventive treatment of migraine. If YES, see the prior authorization guideline related to Migraine Prevention Agents.

Yes No

Q3. The member meets all of the following:

- Is being treated for a diagnosis that is indicated in the U.S. Food and Drug Administration (FDA)-approved package labeling OR a medically accepted indication
- Has a diagnosis confirmed according to the current International Headache Society Classification of Headache Disorders
- Is age-appropriate according to FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature
- Is prescribed a dose that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature
- Does not have a contraindication to the prescribed drug

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Q4. For a gepant for the acute treatment of migraine, one of the following:

- . Has a history of therapeutic failure of at least two (5-HT 1B/1D) receptor agonists (triptans)
- . Has a contraindication or an intolerance to the preferred triptans

If YES, go to 8.

 Yes

 No

 NA

Q5. For a ditran, has a history of therapeutic failure of or a contraindication or an intolerance to the preferred triptans.

If YES, go to 12.

 Yes

 No

 NA

Q6. For an ergot alkaloids, has a history of therapeutic failure of or a contraindication or an intolerance to standard first-line abortive drugs based on headache classification as recommended by current consensus guidelines (such as guidelines from the American Academy of Neurology, American Academy of Family Physicians, American Headache Society).

If YES, go to 12.

 Yes

 No

 NA

Q7. For a non-preferred triptan, has a history of therapeutic failure of or a contraindication or an intolerance to the preferred triptans.

If YES, go to 12.

 Yes

 No

 NA

Q8. For a non-preferred gepant, has a history of therapeutic failure of or a contraindication or an intolerance to the preferred gepants.

If YES, go to 12.

 Yes

 No

 NA



**HEALTH PARTNERS PLANS
PRIOR AUTHORIZATION REQUEST FORM**

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Member Name:

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Q9. For a non-preferred non-steroidal anti-inflammatory drug (NSAID) (e.g., Elyxyb [celecoxib] solution, diclofenac potassium powder packet), has a history of therapeutic failure of or a contraindication or an intolerance to the preferred oral NSAIDs (excluding ketorolac) approved or medically accepted for the beneficiary's diagnosis in the NSAIDs Statewide PDL class.

If YES, go to 12.

Yes

No

NA

Q10. For a non-preferred triptan-NSAID combination product, all of the following:

- . Has a history of therapeutic failure of or a contraindication or an intolerance to the preferred triptans,
- . Has a clinical reason as documented by the prescriber why the individual active ingredients cannot be used concurrently,
- . In addition, for Symbravo (meloxicam-rizatriptan), has a history of therapeutic failure of or a contraindication or an intolerance to sumatriptan-naproxen tablet

If YES, go to 12.

Yes

No

NA

Q11. For non-preferred Migraine Acute Treatment Agents other than triptans, gepants, NSAIDs, and triptan-NSAID combination products (e.g., ditans, ergot alkaloids, etc.), has a history of therapeutic failure of or a contraindication or an intolerance to the preferred Migraine Acute Treatment Agents approved or medically accepted for the beneficiary's diagnosis or indication.

Yes

No

Q12. If a prescription for a Migraine Acute Treatment Agent is for a quantity that exceeds the quantity limit, the determination of whether the prescription is medically necessary will also take into account the guidelines set forth in the plan's quantity limit guidelines. If YES, go to 13.

Yes

No

NA

Q13. The beneficiary is prescribed the requested drug by one of the following:

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Member Name:	Prescriber Name:		
<input type="checkbox"/> A neurologist <input type="checkbox"/> A headache specialist who is certified in headache medicine by the United Council for Neurologic Subspecialties (UCNS)			
<p>Q14. For the acute treatment of migraine, the member is using the requested drug in addition to at least one drug for migraine prevention (e.g., beta-blocker, anticonvulsant, antidepressant, CGRP monoclonal antibody, gepant, botulinum toxin).</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>			
<p>Q15. The member has a history of therapeutic failure of or a contraindication or an intolerance to all preventive migraine drugs recommended by current consensus guidelines (such as guidelines from the American Academy of Neurology, American Academy of Family Physicians, American Headache Society).</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>			
<p>Q16. The member has documentation of an evaluation for the overuse of abortive drugs, including opioids.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>			
<p>Q17. The member meets both of the following:</p> <table> <tr> <td><input type="checkbox"/> Is prescribed a dose that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature</td> <td><input type="checkbox"/> Does not have a contraindication to the prescribed drug</td> </tr> </table>		<input type="checkbox"/> Is prescribed a dose that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature	<input type="checkbox"/> Does not have a contraindication to the prescribed drug
<input type="checkbox"/> Is prescribed a dose that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature	<input type="checkbox"/> Does not have a contraindication to the prescribed drug		
<p>Q18. Documentation is attached showing improvement in headache pain, symptoms, or duration</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>			
<p>Q19. For a non-preferred triptan, has a history of therapeutic failure of or a contraindication or an intolerance to the preferred triptans. If YES, go to 24.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> NA</p>			

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Member Name:	Prescriber Name:
<p>Q20. For a non-preferred gepant, has a history of therapeutic failure of or a contraindication or an intolerance to the preferred gepants. If YES, go to 24.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> NA</p>	
<p>Q21. For a non-preferred NSAID (e.g., Elyxyb [celecoxib] solution, diclofenac potassium powder packet), has a history of therapeutic failure of or a contraindication or an intolerance to the preferred oral NSAIDs (excluding ketorolac) approved or medically accepted for the beneficiary's diagnosis in the NSAIDs Statewide PDL class. If YES, go to 24.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> NA</p>	
<p>Q22. For a non-preferred triptan-NSAID combination product, all of the following:</p> <ul style="list-style-type: none"> . Has a history of therapeutic failure of or a contraindication or an intolerance to the preferred triptans, . Has a clinical reason as documented by the prescriber why the individual active ingredients cannot be used concurrently, . In addition, for Symbravo (meloxicam-rizatriptan), has a history of therapeutic failure of or a contraindication or an intolerance to sumatriptan-naproxen tablet <p>If YES, go to 24.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> NA</p>	
<p>Q23. For non-preferred Migraine Acute Treatment Agents other than triptans, gepants, NSAIDs, and triptan-NSAID combination products (e.g., ditans, ergot alkaloids, etc.), has a history of therapeutic failure of or a contraindication or an intolerance to the preferred Migraine Acute Treatment Agents approved or medically accepted for the beneficiary's diagnosis or indication.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q24. If a prescription for a Migraine Acute Treatment Agent is for a quantity that exceeds the quantity limit, the determination of whether the prescription is medically necessary will also take into account the guidelines set forth in the plan's quantity limit guidelines.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> NA</p>	

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Member Name:	Prescriber Name:
<p>Q25. The beneficiary is prescribed the requested drug by one of the following:</p> <p><input type="checkbox"/> A neurologist <input type="checkbox"/> A headache specialist who is certified in headache medicine by the United Council for Neurologic Subspecialties (UCNS)</p>	
<p>Q26. For the acute treatment of migraine, the member is using the requested drug in addition to at least one drug for migraine prevention (e.g., beta-blocker, anticonvulsant, antidepressant, CGRP monoclonal antibody, gepant, botulinum toxin). If YES, go to 28.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q27. The member has a history of therapeutic failure of or a contraindication or an intolerance to all preventive migraine drugs recommended by current consensus guidelines (such as guidelines from the American Academy of Neurology, American Academy of Family Physicians, American Headache Society).</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q28. The member has documentation of an evaluation for the overuse of abortive drugs, including opioids.</p> <p><input type="checkbox"/> Yes <input type="checkbox"/> No</p>	
<p>Q29. Additional Information:</p>	

Prescriber Signature_____
Date

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