



(such as guidelines from American Academy of Neurology, American Academy of Family Physicians, American Headache Society) (medication, start date and end date): \_\_\_\_\_

- Has documentation of an evaluation for the overuse of abortive medications, including opioids

**SUBMIT MEDICAL RECORD INFORMATION FOR EACH APPLICABLE ITEM.**

**REQUEST FOR GEPANT (NURTEC ODT, UBRELVY) FOR ACUTE TREATMENT OF MIGRAINE:**

- Documented history of therapeutic failure, contraindication or intolerance to at least 2 5-HT<sub>1B/1D</sub> receptor agonists (triptans): (medication, start date and end date): \_\_\_\_\_
- If currently using a different gepant, one of the following:
- Will discontinue use of the other gepant prior to starting the requested gepant
  - Has medical reason for concomitant use of both gepants supported by peer-reviewed literature or national treatment guidelines

**REQUEST FOR DITANS (REYVOW):**

- Documented history of therapeutic failure, contraindication or intolerance to the preferred triptans (medication, start date and end date): \_\_\_\_\_

**REQUEST FOR ERGOT ALKALOIDS (DIHYDROERGOTAMINE):**

- Documented history of therapeutic failure, contraindication or intolerance to standard first-line abortive medications based on headache classification as recommended by current consensus guidelines (such as guidelines from American Academy of Neurology, American Academy of Family Physicians, American Headache Society) (medication, start date and end date): \_\_\_\_\_

**REQUEST FOR NON-PREFERRED MIGRAINE ACUTE TREATMENT:**

- For triptan:
- Documented history of therapeutic failure, contraindication or intolerance to preferred triptans (medication, start date and end date): \_\_\_\_\_
- For all other non-preferred Migraine Acute Treatment:
- Documented history of therapeutic failure, contraindication or intolerance to preferred Migraine Acute Treatment approved or medically accepted for the member's diagnosis (medication, start date and end date): \_\_\_\_\_

**RENEWAL REQUESTS:**

- Member has experienced an improvement in headache pain, symptoms or duration \_\_\_\_\_

**IV. ADDITIONAL RATIONALE FOR REQUEST / PERTINENT CLINICAL INFORMATION :**

Appropriate clinical information to support the request on the basis of medical necessity must be submitted.	Provider Signature:	Date:
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Pharmacy Department will respond via fax or phone within 24 hours.  
Requests for prior authorization (PA) requests must include member name, ID#, and drug name. Please include lab reports with requests when appropriate (e.g., Culture and Sensitivity; Hemoglobin A1C; Serum Creatinine; CD4; Hematocrit; WBC, etc.)