IMPORTANT REMINDER: All providers must revalidate the Medical Assistance (MA) enrollment of each service location every 5 years. Providers should log into PROMISe to check the revalidation dates of each service location and submit revalidation applications at least 60 days prior to the revalidation dates. Enrollment (revalidation) applications may be found at: https://www.dhs.pa.gov/providers/Providers/Pages/PROMISe-Enrollment.aspx.

PURPOSE:

The purpose of this bulletin is to issue updated handbook pages that include the requirements for prior authorization and the type of information needed to evaluate the medical necessity of prescriptions for Migraine Acute Treatment Agents submitted for prior authorization.

SCOPE:

This bulletin applies to all licensed pharmacies and prescribers enrolled in the Medical Assistance (MA) Program. The guidelines to determine the medical necessity of Migraine Acute Treatment Agents will be utilized in the fee-for-service delivery system and by the MA managed care organizations (MCOs) in Physical Health HealthChoices and Community HealthChoices. Providers rendering services in the MA managed care delivery system should address any questions related to the prior authorization of Migraine Acute Treatment Agents to the appropriate MCO.

BACKGROUND:

| *01-21-33 | 09-21-32 | 27-21-24 | 33-21-32 |
| 02-21-20 | 11-21-22 | 30-21-27 |
| 03-21-20 | 14-21-23 | 31-21-35 |
| 08-21-35 | 24-21-30 | 32-21-20 |

COMMENTS AND QUESTIONS REGARDING THIS BULLETIN SHOULD BE DIRECTED TO:

The appropriate toll-free number for your provider type.

Visit the Office of Medical Assistance Programs website at https://www.dhs.pa.gov/providers/Providers/Pages/Health%20Care%20for%20Providers/Contact-Information-for-Providers.aspx.
The Department of Human Services’ (Department) Pharmacy and Therapeutics (P&T) Committee reviews published peer-reviewed medical literature and recommends the following:

- Preferred or non-preferred status for new drugs in therapeutic classes already included in the Preferred Drug List (PDL);
- Changes in the status of drugs on the PDL from preferred to non-preferred and non-preferred to preferred;
- New quantity limits;
- Classes of drugs to be added to or deleted from the PDL; and
- New guidelines or revisions to existing guidelines to evaluate the medical necessity of prescriptions submitted for prior authorization.

DISCUSSION:

During the September 15, 2021, meeting, the P&T Committee recommended the following revisions to the guidelines to determine medical necessity of Migraine Acute Treatment Agents:

- Addition of a guideline that refers to the prior authorization guidelines for Migraine Prevention Agents for requests for a gepant when used for the preventive treatment of migraine;
- Revision of the medical necessity guidelines for gepants to specify which guidelines apply to a gepant prescribed for the acute treatment of migraine;
- Removal of the guideline that a beneficiary will not be using the requested gepant with a calcitonin gene-related peptide monoclonal antibody; and
- Revision of the guideline regarding concomitant use of more than one gepant.

The revisions to the guidelines to determine medical necessity of prescriptions for Migraine Acute Treatment Agents submitted for prior authorization, as recommended by the P&T Committee, were subject to public review and comment and subsequently approved for implementation by the Department.

PROCEDURE:

The procedures for prescribers to request prior authorization of Migraine Acute Treatment Agents are located in SECTION I of the Prior Authorization of Pharmaceutical Services Handbook. The Department will take into account the elements specified in the clinical review guidelines (which are included in the provider handbook pages in the SECTION II chapter related to Migraine Acute Treatment Agents) when reviewing the prior authorization request to determine medical necessity.

As set forth in 55 Pa. Code § 1101.67(a), the procedures described in the handbook pages must be followed to ensure appropriate and timely processing of prior authorization requests for drugs that require prior authorization.
ATTACHMENTS:
Prior Authorization of Pharmaceutical Services Handbook - Updated pages

RESOURCES:
Prior Authorization of Pharmaceutical Services Handbook – SECTION I
Pharmacy Prior Authorization General Requirements
https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/Pharmacy-Prior-Authorization-General-Requirements.aspx

Prior Authorization of Pharmaceutical Services Handbook – SECTION II
Pharmacy Prior Authorization Guidelines
https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/Clinical-Guidelines.aspx
I. Requirements for Prior Authorization of Migraine Acute Treatment Agents

A. Prescriptions That Require Prior Authorization

Prescriptions for Migraine Acute Treatment Agents that meet any of the following conditions must be prior authorized:

1. A prescription for a small molecule calcitonin gene-related peptide (CGRP) receptor antagonist (gepant).

2. A prescription for a serotonin (5-HT) 1F receptor agonist (ditan).

3. A prescription for an ergot alkaloid.

4. A non-preferred Migraine Acute Treatment Agent. See the Preferred Drug List (PDL) for the list of preferred Migraine Acute Treatment Agents at: https://papdl.com/preferred-drug-list.

5. A Migraine Acute Treatment Agent with a prescribed quantity that exceeds the quantity limit. The list of drugs that are subject to quantity limits, with accompanying quantity limits, is available at: https://www.dhs.pa.gov/providers/Pharmacy-Services/Pages/Quantity-Limits-and-Daily-Dose-Limits.aspx.

6. A Migraine Acute Treatment Agent when there is a record of a recent paid claim for another Migraine Acute Treatment Agent in the Point-of-Sale On-Line Claims Adjudication System (therapeutic duplication).

B. Review of Documentation for Medical Necessity

In evaluating a request for prior authorization of a prescription for a Migraine Acute Treatment Agent, the determination of whether the requested prescription is medically necessary will take into account the whether the beneficiary:

1. For a gepant for the preventive treatment of migraine, see the provider handbook pages in the SECTION II chapter related to Migraine Prevention Agents; OR

2. Both of the following:

   a. 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
   b. Has a diagnosis confirmed according to the current International Headache Society Classification of Headache Disorders;

   AND

3. Is age-appropriate according to FDA-approved package labeling, nationally recognized
compendia, or peer-reviewed medical literature; **AND**

4. Is prescribed a dose that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature; **AND**

5. Does not have a contraindication to the prescribed medication; **AND**

6. For a gepant for the acute treatment of migraine, **both** of the following:
   
   a. **One** of the following:
      
      i. Has a history of therapeutic failure of at least two (5-HT\textsubscript{1B/1D}) receptor agonists (triptans)
      ii. Has a contraindication or intolerance to the preferred triptans

   b. If currently using a different gepant, **one** of the following:
      
      i. Will discontinue use of that gepant prior to starting the requested gepant
      ii. Has a medical reason for concomitant use of both gepants that is supported by peer-reviewed medical literature or national treatment guidelines;

   **AND**

7. For a ditan, has a history of trial and failure, contraindication, or intolerance to the preferred triptans; **AND**

8. For ergot alkaloids, has a history of trial and failure, contraindication, or intolerance to standard first-line abortive medications 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); **AND**

9. For a non-preferred Migraine Acute Treatment Agent, **one** of the following:
   
   a. For a non-preferred triptan, has a history of therapeutic failure, contraindication, or intolerance to the preferred triptans
   b. For all other non-preferred Migraine Acute Treatment Agents (e.g., gepants, ditans, ergot alkaloids, etc.), has a history of therapeutic failure, contraindication, or intolerance to the preferred Migraine Acute Treatment Agents approved or medically accepted for the beneficiary’s diagnosis or indication;

   **AND**

10. For therapeutic duplication, **one** of the following:
   
   a. Is being titrated to or tapered from another drug in the same class
b. Has a medical reason for concomitant use of the requested medications that is supported by peer-reviewed literature or national treatment guidelines;

AND

11. 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 all of the following:

a. The guidelines set forth in the Quantity Limits Chapter,

b. Whether the beneficiary is prescribed the requested medication by one of the following:
   
i. A neurologist  
ii. A headache specialist who is certified in headache medicine by the UCNS,

c. For the acute treatment of migraine, both of the following:
   
i. One of the following:
      
      a) The beneficiary is using the requested medication in addition to at least one medication for migraine prevention (e.g., beta-blocker, anticonvulsant, antidepressant, CGRP monoclonal antibody)
      
b) The beneficiary has a history of therapeutic failure, contraindication, or intolerance to all preventive migraine medications recommended by current consensus guidelines (such as guidelines from the American Academy of Neurology, American Academy of Family Physicians, American Headache Society)
   
   ii. Has documentation of an evaluation for the overuse of abortive medications, including opioids.

NOTE: If the beneficiary does not meet the clinical review guidelines listed above but, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary, the request for prior authorization will be approved.

FOR RENEWALS OF PRIOR AUTHORIZATION FOR A MIGRAINE ACUTE TREATMENT AGENT: The determination of medical necessity of a request for renewal of a prior authorization for a Migraine Acute Treatment Agent that was previously approved will take into account whether the beneficiary:

1. Is prescribed a dose that is consistent with FDA-approved package labeling, nationally recognized compendia, or peer-reviewed medical literature; AND
2. Does not have a contraindication to the prescribed medication; **AND**

3. Has documentation of improvement in headache pain, symptoms, or duration; **AND**

4. 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 all of the following:

   a. The guidelines set forth in the Quantity Limits Chapter,

   b. Whether the beneficiary is prescribed the requested medication by one of the following:

      i. A neurologist

      ii. A headache specialist who is certified in headache medicine by the UCNS,

   c. For the acute treatment of migraine, both of the following:

      i. **One** of the following:

         a) The beneficiary is using the requested medication in addition to at least one medication for migraine prevention (e.g., beta-blocker, anticonvulsant, antidepressant, CGRP monoclonal antibody)

         b) The beneficiary has a history of therapeutic failure, contraindication, or intolerance to all preventive migraine medications recommended by current consensus guidelines (such as guidelines from the American Academy of Neurology, American Academy of Family Physicians, American Headache Society)

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

NOTE: If the beneficiary does not meet the clinical review guidelines listed above but, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary, the request for prior authorization will be approved.

B. **Clinical Review Process**

Prior authorization personnel will review the request for prior authorization and apply the clinical guidelines in Section B. above to assess the medical necessity of a prescription for a Migraine Acute Treatment Agent. If the guidelines in Section B. are met, the reviewer will prior authorize the prescription. If the guidelines are not met the prior authorization request will be referred to a physician reviewer for a medical necessity determination. Such a request for prior authorization will be approved when, in the professional judgment of the physician reviewer, the services are medically necessary to meet the medical needs of the beneficiary.

January 3, 2022
(Replacing January 5, 2021)
C. References