

# Pharmacy Drug Policy & Procedure

Policy Name: Reyvow (lasmiditan) Policy #: 2766P

# **Purpose of the Policy**

The purpose of this policy is to establish the criteria for coverage of Reyvow (lasmiditan).

## **Statement of the Policy**

Health Alliance Medical Plans will approve the use of Reyvow (lasmiditan) when the following criteria have been met.

#### Criteria

## 1. Coverage Criteria

- 1.1 Documented diagnosis of moderate to severe migraine, with or without aura, according to the International Classification of Headache Disorders
- 1.2 Age 18 years or older
- 1.3 For patients with 4 or more migraine days per month, documentation that the member is on at least 1 supported migraine prophylactic therapy such as TCAs, SNRIs, beta blockers, anticonvulsants, Botox, etc. with claims history to support adherence through filling at least a 90 day supply within a 120 day time frame
- 1.4 One of the following:
  - Documented trial of at least two generic triptan therapies with little to no relief of moderate/severe migraine symptoms, OR
  - Documented contraindication to triptan therapy defined as one of the following:
    - History of stroke or transient ischemic attack
    - History of hemiplegic or basilar migraine
    - Peripheral vascular disease; ischemic bowel disease
    - Uncontrolled high blood pressure
    - Recent use (within 2 weeks) of MAO inhibitors (e.g., selegiline)
    - Recent use (within 24 hours) of treatment with another 5-HT<sub>1</sub> agonist, or an ergot-containing or ergot-type medication (e.g., methysergide, dihydroergotamine)
    - Ischemic coronary artery disease (angina pectoris, history of myocardial infarction, or documented silent ischemia)
    - Coronary artery vasospasm, including Prinzmetal, variant angina, or other significant underlying cardiovascular disease
    - Wolff-Parkinson-White syndrome or arrhythmias associated with other cardiac accessory conduction pathway disorders
    - Patients with risk factors for CAD (e.g., hypertension, hypercholesterolemia, smoker, obesity, diabetes, strong family history of CAD, menopause, male > 40 years or age) in whom adequate cardiac evaluation has not ruled out CAD

### 2. Managed Dose Limit

- 2.1 Reyvow 50mg: #4 tablets/30 days; Reyvow 100mg: #8 tablets/30 days
- 2.2 The safety of treating an average of more than 4 migraine attacks in a 30 day period has not been established

#### 3. Exclusions

3.1 Reyvow will not be approved if being used in combination with Nurtec ODT (rimegepant) or Ubrelvy (ubrogepant)

## 4. Approval Period

- 4.1 Initial Approval: 12 months
- 4.2 Subsequent Approvals: 12 months with documentation indicating that the member has experienced a positive response to therapy (reduction in pain, photophobia, phonophobia) and claims history indicating that the member has been compliant on migraine prophylactic therapy if the member is having 4 or more migraine headache days per month

CPT Codes	
HCPCS Codes	

#### References

- 1. Olesen J, Bolay H, et al. The International Classification of Headache Disorders, 3<sup>rd</sup> edition. *Cephalagia*. 2018:38(1): 1-211.
- 2. Reyvow (lasmiditan) [prescribing information]. Indianapolis, IN: Lilly USA; September 2022.
- 3. American Headache Society. The American Headache Society Consensus Statement: Update on integrating new migraine treatments into clinical practice. Headache. 2021 Jul;61(7):1021-1039.

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