

# Commercial/Healthcare Exchange PA Criteria

Effective: March 2008

**Prior Authorization:** Eletriptan

**Products Affected:** Relpax (eletriptan) oral tablets, Eletriptan oral tablets

#### **Medication Description:**

Eletriptan has high affinity for 5-HT1 receptor subtypes, more specifically those receptors located on intracranial blood vessels, and acts as a receptor agonist at those sites. Two theories exist to explain the 5-HT1 receptor agonists in migraine headaches: activation of 5-HT1 vasoconstricts intracranial blood vessels and correlates to relief of migraine headaches; activation of 5-HT1 receptors on sensory nerve endings inhibits pro-inflammatory response in the trigeminal system.

Covered Uses: Eletriptan is indicated for the acute treatment of migraine with or without aura in adults.

#### **Exclusion Criteria:**

- 1. Prevention of migraine attacks
- 2. Cluster headache
- 3. Ischemic coronary artery disease (CAD) (angina pectoris, history of myocardial infarction, or documented silent ischemia) or coronary artery vasospasm, including Prinzmetal's angina
- 4. Wolff-Parkinson-White syndrome or arrhythmias associated with other cardiac accessory conduction pathway disorders
- 5. History of stroke, transient ischemic attack (TIA), or history or current evidence of hemiplegic or basilar migraine because these patients are at a higher risk of stroke
- 6. Peripheral vascular disease
- 7. Ischemic bowel disease
- 8. Uncontrolled hypertension
- 9. Use within 24 hours of treatment with another 5-HT1 agonist, or an ergotamine-containing medication
- 10. Use within at least 72 hours of treatment with the following potent CYP3A4 inhibitors: ketoconazole, itraconazole, nefazodone, troleandomycin, clarithromycin, ritonavir, or nelfinavir

### **Required Medical Information:**

- 1. Diagnosis
- 2. Previous therapies tried and failed

Age Restrictions: 18 years of age and older

**Prescriber Restrictions:** N/A

**Coverage Duration:** 12 months

Other Criteria:



Last Res.6.5.2020



1. Patient has a documented intolerance to, or treatment failure of an adequate trial of TWO of the following medications: naratriptan, almotriptan, frovatriptan, sumatriptan, rizatriptan, or zolmitriptan

## References:

- 1. Product Information: RELPAX(R) oral tablets, eletriptan hydrobromide oral tablets. Roerig (per FDA), New York, NY, 2013.
- 2. Product Information: Eletriptan hydrobromide oral tablets, eletriptan hydrobromide oral tablets. Amneal Pharmaceuticals LLC (per DailyMed), Bridgewater, NJ, 2018.

## **Policy Revision history**

Rev#	Type of Change	Summary of Change	Sections Affected	Date
1	New Policy	All	All	3/2008
2	Template update	Reformatted from old to new CCI Template  CCI P&T Review History: 3/08, 6/08, 9/09, 9/10, 12/11, 10/12, 10/13, 10/14, 11/15, 5/16, 11/16, 5/17, 4/17, 5/18, 5/19  CCI Revision Record: 8/11, 6/13, 8/13, 4/16, 11/16, 5/18  Added Mechanism of Action  Updated Exclusion Criteria to match FDA Label  Updated Reference	All	6/5/2020



3	Update	Updated Exclusion Criteria to match FDA Label with the addition of #9 and #10	Exclusion Criteria	2/10/2021
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