

Commercial/Healthcare Exchange PA Criteria

Effective: September 2010

Prior Authorization: Sumavel DosePro

Products Affected: Sumavel DosePro (sumatriptan) needle-free subcutaneous injection

<u>Medication Description</u>: Sumavel DosePro is a prefilled, single-dose, needle-free subcutaneous delivery system delivering 0.5 mL of sterile solution containing 4 mg or 6 mg sumatriptan (as the succinate salt). Sumavel DosePro contains sumatriptan succinate, a selective 5-HT1B/1D receptor agonist indicated in adults for (1) the acute treatment of migraine, with or without aura, and (2) the acute treatment of cluster headache.

Covered Uses:

- 1. Acute treatment of migraine, with or without aura
- 2. Acute treatment of cluster headache

Exclusion Criteria:

- 1. Prevention of migraine attacks
- 2. Patients with ischemic coronary artery disease (CAD) (angina pectoris, history of myocardial infarction, or documented silent ischemia) or coronary artery vasospasm, including Prinzmetal's angina
- 3. Patients with Wolff-Parkinson-White syndrome or arrhythmias associated with other cardiac accessory conduction pathway disorders
- 4. Patients with a history of stroke or transient ischemic attack (TIA) because these patients are at a higher risk of stroke
- 5. Patients with a history of hemiplegic or basilar migraine
- 6. Patients with peripheral vascular disease
- 7. Patients with ischemic bowel disease
- 8. Patients with uncontrolled hypertension
- 9. Patients with recent (i.e., within 24 hours) use of ergotamine-containing medication, ergot-type medication (such as dihydroergotamine or methysergide), or another 5-hydroxytryptamine1 (5-HT1) agonist
- 10. Concurrent administration of a monoamine oxidase (MAO)-A inhibitor or recent (within 2 weeks) use of an MAO-A inhibitor

Required Medical Information:

- 1. Diagnosis
- 2. Past medication trials

Age Restrictions: 18 years of age and older

<u>Prescriber Restrictions</u>: Prescribed by, or in consultation with, a neurologist or pain management specialist.

Coverage Duration: 12 months

Other Criteria:

- 1. Patient has a documented diagnosis of migraine or cluster headache; AND
- 2. Patient experiences at least two headaches per month; AND

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3. Patient has a documented intolerance, contraindication, or treatment failure with, an adequate trial of an oral triptan (sumatriptan, frovatriptan, rizatriptan, etc.) AND generic sumatriptan subcutaneous injection.

*ConnectiCare does not consider needle-phobia to be a clinical reason to use Sumavel DosePro over injectable medications.

References:

1. Product Information: Sumavel(R) DosePro(R) subcutaneous injection solution, sumatriptan succinate subcutaneous injection solution. Zogenix, Inc. (per FDA), San Diego, CA, 2013.

Policy Revision history

Rev#	Type of Change	Summary of Change	Sections Affected	Date
1	Policy Update	CCI adoption of EH template and criteria. CCI P&T Review History: 9/10, 12/11, 4/12, 10/12, 10/13, 10/14, 11/15, 5/16, 8/16, 11/16, 2/17, 1/18 CCI Revision Record: 4/12, 8/16, 11/16	All	10/15/2019

