

Commercial/Healthcare Exchange PA Criteria Effective: December 2011

Prior Authorization: Gralise

Products Affected: Gralise (gabapentin) oral tablet, Gralise (gabapentin) ER tablets

<u>Medication Description</u>: Gralise is an analog of the neurotransmitter gammaaminobutyric acid (GABA). Gralise exerts its pharmacologic action by binding to the alpha-2-delta subunit of voltage-gated calcium channels. The binding of this subunit reduces the release of several neurotransmitters including glutamate, noradrenaline, and substance P.

Covered Uses: Management of postherpetic neuralgia.

Exclusion Criteria:

1. Known hypersensitivity to gabapentin

Required Medical Information:

1. Diagnosis

2. History of previous medications tried/failed

Age Restrictions: 18 years of age and older

Prescriber Restrictions: N/A

Coverage Duration: 12 months

Other Criteria:

- A. Patient has a diagnosis of postherpetic neuralgia; AND
- B. Patient has had treatment failure, intolerance, or contraindication to tricyclic antidepressants; AND
- C. Patient has had an intolerance to, or treatment failure of, gabapentin tablets or capsules, at a minimum dose of 1800mg per day.

References:

- 1. Gralise Full Prescribing Information, Menlo Park, CA, Depomed Inc
- 2. Facts & Comparisons online

Policy Revision history

Rev#	Type of Change	Summary of Change	Sections Affected	Date
1	New Policy	New Policy	All	12/2011
2.	Update	Moved to updated template CCI Revision Record: 9/15, 11/17	All	2/3/2020

Last Rev. May 19,.2023





3	Update	Added Gralise ER tablets to Products Affected	Products Affected	5/19/2023	
---	--------	--	-------------------	-----------	--