



For the acute treatment of migraine with or without aura  
and the preventive treatment of episodic migraine in adults

**Nurtec**<sup>®</sup> ODT  
(rimegepant)  
orally disintegrating tablets 75 mg

## American Headache Society Goals for Acute Treatment of Migraine

**The 2021 American Headache Society (AHS) Consensus Statement recommends 6 goals for acute treatment of migraine<sup>1</sup>**



Rapid and consistent freedom from pain and associated symptoms, especially the most bothersome symptom, without recurrence



Optimal self-care and reduced subsequent use of resources (eg, emergency room visits, diagnostic imaging, clinician and ambulatory infusion center visits)



Minimal need for repeat dosing or rescue medications



Restored ability to function



Minimal or no adverse events



Cost considerations

**Suboptimal acute treatment is associated with higher migraine-related disability and risk of disease progression<sup>1</sup>**

**Consider Nurtec ODT as an acute treatment option for your patients with migraine**

CGRP=calcitonin gene-related peptide.

\*Per IQVIA as oral brand in class (oral CGRP receptor antagonists): number one prescribed and number one in new prescriptions, since 8/6/21. Data current as of 10/30/24.

### SELECT IMPORTANT SAFETY INFORMATION

**Contraindications:** Hypersensitivity to Nurtec ODT or any of its components.

### Warnings and Precautions

**Hypersensitivity Reactions:** If a serious hypersensitivity reaction occurs, discontinue Nurtec ODT and initiate appropriate therapy. Serious hypersensitivity reactions have included dyspnea and rash and can occur days after administration.

**Please see additional Important Safety Information on the next page and click here for full [Prescribing Information](#).**



**Learn how Nurtec ODT may help your patients with migraine.**

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## INDICATIONS

Nurtec ODT is indicated in adults for the:

- acute treatment of migraine with or without aura
- preventive treatment of episodic migraine

## IMPORTANT SAFETY INFORMATION

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### Warnings and Precautions

**Hypersensitivity Reactions:** If a serious hypersensitivity reaction occurs, discontinue Nurtec ODT and initiate appropriate therapy. Serious hypersensitivity reactions have included dyspnea and rash and can occur days after administration.

**Hypertension:** Development of hypertension and worsening of pre-existing hypertension have been reported following the use of CGRP antagonists, including Nurtec ODT, in the postmarketing setting. Monitor patients for new-onset hypertension or worsening of pre-existing hypertension and consider whether discontinuation is warranted.

**Raynaud's Phenomenon:** Development of Raynaud's phenomenon and recurrence or worsening of pre-existing Raynaud's phenomenon have been reported in the postmarketing setting following the use of CGRP antagonists, including Nurtec ODT.

If signs or symptoms of Raynaud's phenomenon develop, discontinue Nurtec ODT. Patients should be evaluated by a healthcare provider if symptoms do not resolve. Patients with a history of Raynaud's phenomenon should be monitored for and informed about the possibility of worsening or recurrence of signs and symptoms.

**Adverse Reactions:** The most common adverse reactions for Nurtec ODT vs placebo were nausea (2.7% vs 0.8%) and abdominal pain/dyspepsia (2.4% vs 0.8%).

**Drug Interactions:** Avoid concomitant administration of Nurtec ODT with strong inhibitors of CYP3A4 or strong or moderate inducers of CYP3A. Avoid another dose of Nurtec ODT within 48 hours when it is administered with moderate inhibitors of CYP3A4 or potent inhibitors of P-gp.

**Use in Specific Populations:** *Pregnancy:* It is not known if Nurtec ODT can harm an unborn baby. *Lactation:* The transfer of rimegepant into breast milk is low (<1%). *Hepatic impairment:* Avoid use of Nurtec ODT in persons with severe hepatic impairment. *Renal impairment:* Avoid use in patients with end-stage renal disease.

**Please click here for full [Prescribing Information](#).**

**Reference: 1.** Ailani J, Burch RC, Robbins MS; the Board of Directors of the American Headache Society. The American Headache Society Consensus Statement: Update on integrating new migraine treatments into clinical practice. *Headache*. 2021;61:1021-1039.