

**For the acute treatment of migraine in adults.
Not for preventive treatment.**

Nurtec™ ODT
(rimegepant)
orally disintegrating tablets 75 mg

Getting Started Prescribing Nurtec™ ODT (rimegepant)

Just one dose can:¹



Dissolve Pain

Quick-dissolve tablet can start working in minutes to eliminate pain¹⁻³



Restore Function

Proven to get many patients back to normal activities in as little as 1 hour^{1,2,4}



Last up to 48 hours

Can keep patients functioning and free from migraine pain for up to 2 days—without rescue medication^{1,2,5}

In a multi-center, double-blind, randomized, placebo-controlled study of 1351 patients (Nurtec ODT 75 mg, n=669; placebo, n=682), co-primary endpoints at 2 h for Nurtec ODT vs placebo were: pain freedom (21% vs 11%, $P < .001$) and freedom from most bothersome symptom (MBS; predefined as photophobia, phonophobia, or nausea; 35% vs 27%, $P = .001$). Select secondary endpoints at 60 minutes for Nurtec ODT vs placebo: pain relief (37% vs 31%, $P < .05$) and return to normal function (22% vs 16%, $P = .0025$); from 2-48 h for Nurtec ODT vs placebo: pain freedom (14% vs 5%, $P < .001$), and return to normal function (26% vs 15%, $P < .0001$).^{1,2,4,5}

INDICATION

Nurtec ODT is indicated for the acute treatment of migraine with or without aura in adults.

Limitations of Use

Nurtec ODT is not indicated for the preventive treatment of migraine.

SELECT IMPORTANT SAFETY INFORMATION

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

Please see additional Important Safety Information on last page and the full Prescribing Information.



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Getting started with Nurtec™ ODT (rimegepant)



Prescribing Nurtec ODT 75 mg is simple¹

Each 8-pack of Nurtec ODT is meant to cover 8 migraine attacks, as needed up to once daily.¹

Prescription: Once daily PRN for migraine.

The safety of treating more than 15 migraines in a 30-day period has not been established.¹

The Nurtec ODT Savings Card

An activated copay card is included in the Patient Starter Kit. Patients can also download and activate a card to get started today via 2 easy ways:

- visit www.Nurtec.com/savings
- text NSAVE to 267-89[†]

Available for eligible patients with commercial insurance.

See last page for Terms & Conditions.



**Patients can pay as little as
\$0 per month***

*Per 8-tablet prescription. See Terms & Conditions.

[†]Terms and conditions apply. Message and data rates may apply. Patients may receive approximately 5 messages a month. For information, including our Privacy Policy and Terms & Conditions, please visit www.Nurtec.com/copay-terms. Once registered, text HELP to 267-89 for more information. Text STOP to end.



We're here to help.

Call **1-833-4NURTEC** (M-F from 8 am - 8 pm ET) or visit www.Nurtec-HCP.com/savings-support to get your patients started today!

SELECT IMPORTANT SAFETY INFORMATION

Warnings and Precautions: 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 last page and the full Prescribing Information.

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Helpful resources for you and your patients

- 8-tablet packs of Nurtec ODT for treating up to 8 migraines
- Formulary Coverage tool to obtain up-to-date coverage in your area (by zip code)
- Prior Authorization assistance
- Copay Card
- Biohaven Patient Assistance Program



Prior Authorization (PA) in three easy steps:

- 1 Log in**
Log in to your existing CoverMyMeds account or create one at covermymeds.com.
- 2 Complete a request**
Generate a PA request required for treatment, or complete a pharmacy-initiated request.
- 3 Electronically submit**
Click one button to submit the request to patient's plan for determination.

Prior Authorization must be completed to prevent disruption to your patient's therapy.

Need help getting started?

Contact CoverMyMeds: **1-866-452-5017** | go.covermymeds.com/help

For prescribing/PA support resources, visit: <https://www.nurtec-hcp.com/resources>

SELECT IMPORTANT SAFETY INFORMATION

Adverse Reactions: The most common adverse reaction was nausea (2% in patients who received Nurtec ODT compared to 0.4% in patients who received placebo). Hypersensitivity, including dyspnea and rash, occurred in less than 1% of patients treated with Nurtec ODT.

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Savings Program Terms & Conditions

To the Patient: In order to participate in the Nurtec ODT Patient Savings Program ("Program"), you must have a valid prescription for Nurtec™ ODT (rimegepant) orally disintegrating tablets, meet the eligibility requirements set forth herein, adhere to the terms and conditions stated in the Restrictions section below, and present this card to your pharmacist. Eligible patients with commercial insurance may pay as little as \$0 out of pocket and limited to one 8-tablet prescription per month. Patients with questions about the Program should call 1-800-761-1568.

To the Pharmacist: By redeeming this offer, the Pharmacist certifies: (a) that the Pharmacy has not submitted, and will not submit, a claim for reimbursement under any federal, state, or other government programs for this prescription or where prohibited by law and (b) the Pharmacist will adhere to the terms and conditions stated in the Restrictions section below.

Pharmacist Instructions: For Commercially Insured Patients, please submit this claim to the patient's primary Third Party Payer first, then submit the balance due to **CHANGE HEALTHCARE** as a Secondary Payer COB (coordination of benefits) with patient responsibility and a valid Other Coverage Code (e.g. 08). Reimbursement will be received from **CHANGE HEALTHCARE**. For questions, please call the Concierge line for Nurtec ODT at 1-800-731-4997, Monday – Friday, 8 am – 8 pm ET.

Restrictions: This offer is not valid for Non-Insured/Cash-Paying Patients. This Program is not valid for prescriptions covered by or submitted for reimbursement in part or in full by any state or federally funded programs, including but not limited to Medicare, Medicaid, Medigap, VA, TRICARE (DOD). Patients with managed care restrictions (e.g., prior authorization, step edit) may not be eligible for this offer if such managed care restrictions persist. Continued eligibility may require that the patient has a prior authorization form submitted. This offer may not be used with any other financial assistance program, free trial, discount, prescription savings card or other offer. Valid only for patients 18 years and older in the United States including the Commonwealth of Puerto Rico. This Program is void if copied, transferred, purchased, altered, or traded and where prohibited by law. Limit one offer per individual. This offer expires on December 31, 2021. Biohaven Pharmaceuticals reserves the right to rescind, revoke or amend this offer without notice anytime. This Program is managed by ConnectiveRx on behalf of Biohaven Pharmaceuticals.

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IMPORTANT SAFETY INFORMATION

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Adverse Reactions: The most common adverse reaction was nausea (2% in patients who received Nurtec ODT compared to 0.4% in patients who received placebo). Hypersensitivity, including dyspnea and rash, occurred in less than 1% of patients treated with Nurtec ODT.

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

Use in Specific Populations: *Pregnant/breast feeding:* It is not known if Nurtec ODT can harm an unborn baby or if it passes into breast milk. *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 see full [Prescribing Information](#).

Price disclosure information for prescribers available here: [Nurtec-HCP.com/pricing](https://www.biohaven.com/pricing)

REFERENCES: **1.** Nurtec ODT. Package insert. Biohaven Pharmaceuticals Inc. **2.** Croop R, Goadsby PJ, Stock DA, et al. Efficacy, safety, and tolerability of rimegepant orally disintegrating tablet for the acute treatment of migraine: a randomised, phase 3, double-blind, placebo-controlled trial. *Lancet*. 2019;394(10200):737-745. doi: 10.1016/S0140-6736(19)31606-X.

3. Data on File. RIM108. Biohaven Pharmaceuticals Inc.

4. Data on File. RIM107. Biohaven Pharmaceuticals Inc.

5. Data on File. RIM118. Biohaven Pharmaceuticals Inc.

