

UnitedHealthcare Pharmacy
Clinical Pharmacy Programs

Program Number	2022 P 1326-4
Program	Prior Authorization-Notification
Medication	Nurtec ODT (rimegepant), Qulipta (atogepant)*, Ubrelvy (ubrogepant)
P&T Approval Date	9/2020, 7/2021, 12/2021, 3/2022
Effective Date	4/1/2022; Oxford only: N/A

1. Background:

Nurtec ODT (rimegepant) and Ubrelvy (ubrogepant) are calcitonin gene-related peptide receptor antagonists indicated for the acute treatment of migraine with or without aura in adults. Nurtec ODT and Qulipta (atogepant)* are indicated for the preventive treatment of episodic migraine in adults.

2. Coverage Criteria^a:

A. Initial Authorization

1. **Ubrelvy** will be approved based on **all** of the following criteria:

a. Diagnosis of migraine headaches with or without aura.

-AND-

b. Used for acute treatment of migraine

-AND-

c. Medication will not be used in combination with another acute calcitonin gene-related peptide receptor (CGRP) antagonist (i.e., Nurtec ODT)

Authorization will be issued for 12 months.

2. Nurtec ODT will be approved based on **one** of the following criteria:

a. **All** of the following:

1) Diagnosis of migraine headaches with or without aura.

-AND-

2) Used for acute treatment of migraine

-AND-

3) Medication will not be used in combination with another acute calcitonin gene-related peptide receptor (CGRP) antagonists (i.e., Ubrelvy)

-OR-

b. **All** of the following:

1) Diagnosis of episodic migraines

-AND-

2) Used for preventive treatment of migraines

-AND-

3) Medication will not be used in combination with another CGRP antagonist or inhibitor used for the preventive treatment of migraines (e.g., Aimovig, Ajovy*, Emgality, Qulipta*, Vyepti^)

Authorization will be issued for 12 months.

3. Qulipta* will be approved based all of the following criteria:

a. Diagnosis of episodic migraines

-AND-

b. Used for preventive treatment of migraines

-AND-

c. Medication will not be used in combination with another CGRP antagonist or inhibitor used for the preventive treatment of migraines (e.g., Aimovig, Ajovy*, Emgality, Vyepti^, Nurtec ODT)

Authorization will be issued for 12 months.

B. Reauthorization

1. **Ubrelvy** will be approved based on the following criteria:

a. Documentation of positive clinical response to therapy

-AND-

b. Medication will not be used in combination with another acute calcitonin gene-related peptide receptor (CGRP) antagonist (i.e., Nurtec ODT)

Authorization will be issued for 12 months.

2. **Nurtec ODT** will be approved based on the following criteria:

- a. Documentation of positive clinical response to therapy

-AND-

- b. **One** of the following:

- 1) **Both** of the following:

- a) Use is for the acute treatment of migraine

-AND-

- b) Medication will not be used in combination with another acute calcitonin gene-related peptide receptor (CGRP) antagonist (i.e., Ubrelvy)

-OR-

- 2) **Both** of the following:

- a. Use is for the preventive treatment of migraines

-AND-

- b. Medication will not be used in combination with another CGRP antagonist or inhibitor used for the preventive treatment of migraines (e.g., Aimovig, Ajovy*, Emgality, Qulipta*, Vyepti^)

Authorization will be issued for 12 months.

3. **Qulipta*** will be approved based on the following criteria:

- a. Documentation of positive clinical response to therapy

-AND-

- b. Medication will not be used in combination with another CGRP antagonist or inhibitor used for the preventive treatment of migraines (e.g. Aimovig, Ajovy*, Emgality, Nurtec ODT, Vyepti^)

Authorization will be issued for 12 months.

^a State mandates may apply. Any federal regulatory requirements and the member specific benefit plan coverage may also impact coverage criteria. Other policies and utilization management programs may apply.

Ajovy and Qulipta are typically excluded from coverage

^Vyepti may be subject to additional benefit and coverage review requirements.

3. Additional Clinical Programs:

- Supply limits may apply.
- Prior Authorization-Medical Necessity may apply
- Step Therapy may apply
- Notwithstanding Coverage Criteria, UnitedHealthcare may approve initial and re-authorization based solely on previous claim/medication history, diagnosis codes (ICD-10) and/or claim logic. Use of automated approval and re-approval processes varies by program and/or therapeutic class.

4. References:

1. Nurtec ODT [package insert]. New Haven, CT: Biohaven Pharmaceuticals, Inc.; December 2021.
2. Qulipta [package insert]. Dublin, Ireland: Forest Laboratories Ireland, Ltd. October 2021.
3. Ubrelvy [package insert]. Madison, NJ: Allergan USA, Inc.; March 2021.
4. The American Headache Society Position Statement on Integrating New Migraine Treatments Into Clinical Practice. AHS Consensus Statement. Headache. 2021; 61:1021-39.

Program	Prior Authorization-Notification – Nurtec ODT, Ubrelvy
Change Control	
Date	Change
9/2020	New program.
7/2021	Added new indication for Nurtec ODT for episodic migraines. Added statement regarding concomitant therapy with other CGRPs. Updated references. Added state mandate language.
12/2021	Added Qulipta for preventive treatment of migraines. Updated references.
3/2022	Updated the products typically excluded from coverage. Added note for Vyetpi regarding additional benefit and coverage review requirements. Updated references.