Norliqva®
(Amlodipine) Oral Solution
1 mg/mL

# Only Norliqva checks all of these boxes[1,2]




| | Norliqva® (amlodipine) Oral Solution[1,‡] | Katerzia® (amlodipine) Oral Suspension[2,§] |
|---|---|---|
| Alcohol-free | ✓ | ✓ |
| No shaking | ✓ | ✗ |
| No refrigeration | ✓ | ✗ |
| Lower wholesale acquisition costs (WAC)* | ✓ | ✗ |
| Patient copay as low as $0† | ✓ | ✗ |

**This comparison is based on each product's Prescribing Information and published copay program terms (10/13/26). This is not intended to imply comparable indications, efficacy, safety, or clinical outcomes. No head-to-head studies have been conducted. See the full Prescribing Information for each product. Report adverse events or medical information requests for other products to their respective manufacturers.**

## PRODUCT AND ORDERING INFORMATION

### DOSING AND ADMINISTRATION[1]

| | |
|---|---|
| **Norliqva is an oral solution** | **1 mg/mL** |
| **Recommended adult starting dose** | **5 mg orally once daily** with a maximum of **10 mg orally once daily** |
| **Pediatric (ages 6 and older) starting dose** | **2.5 mg** to **5 mg orally once daily** |

Small, fragile, or elderly patients or patients with hepatic insufficiency may be started on 2.5 mg orally once daily. Doses in excess of 5 mg daily have not been studied in pediatric patients. Dispense in light-resistant and tight (per USP) container.

### PRODUCT INFORMATION[1,3]

| | |
|---|---|
| **Bottle** | **150 mL** |
| **Storage** | Stable 24-month shelf life, no refrigeration required |
| **NDC** | **46287-037-01** |
| **Ordering information** | Norliqva can be ordered from your wholesale distributor nationwide |

### WHOLESALER INFORMATION

| | Bottle Size/NDC | Item Number |
|---|---|---|
| **Cardinal** | 150 mL: 46287-037-01 | 6125298 |
| **Cencora** | 150 mL: 46287-037-01 | 10309445 |
| **McKesson** | 150 mL: 46287-037-01 | 3090990 |
| **Morris and Dickson** | 150 mL: 46287-037-01 | 924084 |

## IMPORTANT SAFETY INFORMATION

### Indications and Usage

NORLIQVA is a calcium channel blocker for the treatment of:

**HYPERTENSION**

NORLIQVA is indicated for the treatment of hypertension in adults and children 6 years of age and older, to lower blood pressure. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular events, primarily strokes and myocardial infarctions.

**CORONARY ARTERY DISEASE**

- Chronic Stable Angina
- Vasospastic Angina (Prinzmetal's or Variant Angina)
- Angiographically Documented Coronary Artery Disease In patients without heart failure or an ejection fraction <40%

**Please see full Prescribing Information for additional safety information.**

*Representations herein are based on wholesale acquisition cost (WAC) as published in Medi-Span as of 10/16/26: Norliqva 150-mL bottle $515; Katerzia 150-mL bottle $658.97. WAC is the manufacturer's published list price to wholesalers and does not include discounts, rebates, or other price concessions. WAC does not represent the price paid by pharmacies, payers, or patients. Actual patient cost depends on insurance coverage, formulary status, pharmacy, and other factors. Pricing is subject to change.

†Eligible patients may pay as little as $0 per prescription through the CMP EasyPay Copay Program. Available to commercially insured, commercially uninsured and cash patients. Not valid for patients enrolled in Medicare, Medicaid, TRICARE, VA, or any other state or federally funded healthcare program, or where prohibited by law. Maximum savings of up to $150 off per 1-89-day supply and up to $450 off per 90+ day supply. Patient out-of-pocket cost will vary by insurance plan; patients with high deductibles or non-covered status may pay more than $0. Program is not health insurance, cannot be combined with other offers, and may be modified or discontinued without notice. Full terms and conditions at easypaysavings.com/terms.

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§Katerzia® is a registered trademark of Azurity Pharmaceuticals, Inc. CMP Pharma, Inc. is not affiliated with, sponsored by, or endorsed by Azurity Pharmaceuticals, Inc.

# FDA-approved formulation. No additional preparation.

- **Norliqva is a ready-to-use oral solution that serves as an alternative to pharmacy compounding[1,3,4]**
  - Consistent with AAP guidance recommending use of commercially available, FDA-approved formulations when possible[5,6]
  - Under FDA guidance, pharmacies generally should not compound products that are essentially copies of commercially available, FDA-approved products[7]
  - Pharmacies creating compounded versions of FDA-approved medications may violate the Federal Food, Drug, and Cosmetic Act (FDCA)[7]




**Patients may pay as little as $0 with the EasyPay Copay Program***

**Visit cmppharma.com/easypay for more information**

Restrictions apply, see terms

**Eligible commercially insured and cash-paying patients. Not valid for government-insured patients. Maximum benefit limits apply.**

## IMPORTANT SAFETY INFORMATION (continued)

### Contraindications

NORLIQVA is contraindicated in patients with sensitivity to amlodipine.

### Warnings and Precautions

NORLIQVA may cause the following conditions.

- Symptomatic hypotension is possible, particularly in patients with severe aortic stenosis. However, acute hypotension is unlikely.
- Worsening angina and acute myocardial infarction can develop after starting or increasing the dose of NORLIQVA, particularly in patients with severe obstructive coronary artery disease.
- Titrate slowly in patients with severe hepatic impairment.

### Adverse Reactions

Most common adverse reactions to amlodipine were edema, dizziness, flushing and palpitation which occurred in a dose related manner. Other adverse reactions not clearly dose-related but reported with an incidence >1.0% are fatigue and nausea.

See full Prescribing Information for other possible side effects with NORLIQVA. To report SUSPECTED ADVERSE REACTIONS, contact CMP Pharma, Inc. at 1-844-321-1443 or FDA at 1-800-FDA-1088 or www.fda.gov/medwatch.

### Drug Interactions

Do not exceed doses greater than 20 mg daily of simvastatin.

- CYP3A Inhibitors: Co-administration with CYP3A inhibitors (moderate and strong) results in increased systemic exposure to amlodipine and may require dose reduction.
- CYP3A Inducers: No information is available on the quantitative effects of CYP3A inducers on amlodipine. Blood pressure should be closely monitored when amlodipine is coadministered with CYP3A inducers.
- Simvastatin: Co-administration of simvastatin with amlodipine increases the systemic exposure of simvastatin. Limit the dose of simvastatin in patients on amlodipine to 20 mg daily.
- Immunosuppressants: Amlodipine may increase the systemic exposure of cyclosporine or tacrolimus when co-administered.

### Dosage and Administration

NORLIQVA is an oral solution: 1 mg/mL. Adult recommended starting dose: 5 mg orally once daily with a maximum of 10 mg orally once daily. Small, fragile, or elderly patients or patients with hepatic insufficiency may be started on 2.5 mg orally once daily. Pediatric starting dose: 2.5 mg to 5 mg orally once daily.

**Please see full Prescribing Information for additional safety information.**

**References: 1.** Norliqva. Prescribing information. CMP Pharma, Inc; 2026. **2.** Katerzia. Prescribing information. Azurity Pharmaceuticals, Inc; 2025. **3.** Stockbridge N. *NDA Approval: NDA 214439, Norliqva (Amlodipine) Oral Solution.* US Food and Drug Administration; 2022. **4.** Gudeman J, Jozwiakowski M, Chollet J, et al. Potential risks of pharmacy compounding. *Drugs R D.* 2013;13(1):1-8. doi:10.1007/s40268-013-0005-9 **5.** Flynn JT, Kaelber DC, Baker-Smith CM, et al. Clinical practice guideline for screening and management of high blood pressure in children and adolescents. *Pediatrics.* 2017;140(3):e20171904. doi:10.1542/peds.2017-1904 **6.** Current good manufacturing practice for finished pharmaceuticals. 21 CFR pt 211 (2026). Accessed October 29, 2026. https://www.ecfr.gov/current/title-21/chapter-I/subchapter-C/part-211 **7.** US Food and Drug Administration, Center for Drug Evaluation and Research. *Pharmacy Compounding of Human Drug Products Under Section 503A of the Federal Food, Drug, and Cosmetic Act: Guidance.* Revision 2. US Dept of Health and Human Services; June 2016. Accessed October 21, 2026. https://www.fda.gov/media/94393/download

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