OXYMETAZOLINE HYDROCHLORIDE- oxymetazoline hydrochloride spray
Bi-Mart

Drug Facts

Active ingredient

Oxymetazoline hydrochloride 0.05%

Purpose

Nasal Decongestant

Uses

- temporarily relieves nasal congestion due to:
  - common cold
  - hay fever
  - upper respiratory allergies
- temporarily relieves sinus congestion and pressure
- shrinks swollen nasal membranes so you can breathe more freely

Warnings

Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland

When using this product

- **do not use more than directed**
- do not use for more than 3 days. Use only as directed. Frequent or prolonged use may cause nasal congestion to recur or worsen.
- temporary discomfort such as burning, stinging, sneezing or an increase in nasal discharge may occur
- use of this container by more than one person may spread infection.

Stop use and ask a doctor if

symptoms persist

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away at 1-800-222-1222.

Directions

- **adults and children 6 to under 12 years of age (with adult supervision):**2 or 3 sprays in each nostril not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24-hour period.
- **children under 6 years of age:**ask a doctor.

To use: Push firmly down on cap and turn counter clockwise. **To spray**, squeeze bottle quickly and firmly. Do not tilt head backward while spraying. Wipe nozzle clean after use. Secure cap after use.

Other information

- store at room temperature

Inactive ingredients

benzalkonium chloride, benzyl alcohol, edetate disodium, polyethylene glycol, povidone, propylene glycol, purified water, sodium phosphate dibasic, sodium phosphate monobasic

Questions?

1-866-467-2748

PRINCIPAL DISPLAY PANEL

OXYMETAZOLINE HYDROCHLORIDE

oxymetazoline hydrochloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37835-996
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY)	OXYMETAZOLINE	0.05 g

(OXYMETAZOLINE - UNII:8VLN5B44ZY)	HYDROCHLORIDE	in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37835-996-01	1 in 1 CARTON	08/01/2025	
1		30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	08/01/2025	

Labeler - Bi-Mart (027630078)

Registrant - Bi-Mart (027630078)

Establishment

Name	Address	ID/FEI	Business Operations
RARITAN PHARMACEUTICALS INCORPORATED		127602287	manufacture(37835-996)

Revised: 8/2025 Bi-Mart