

OXYMETAZOLINE HYDROCHLORIDE- oxymetazoline hydrochloride spray
Prodose, Inc.

Oxymetazoline Hydrochloride 0.05% Nasal Spray

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.

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.

Shake well before use. Before using the first time, remove the protective cap from the tip and prime metered pump by depressing pump firmly several times. To spray, hold bottle with thumb at base and nozzle between first and second fingers. Without tilting head, insert nozzle into nostril. Fully depress rim with a firm, even stroke and sniff deeply. Wipe nozzle clean after use.

Other information

- store between 20° to 25°C (68° to 77°F)
- retain carton for future reference on full labelling

Inactive ingredients

benzalkonium chloride solution, benzyl alcohol, edetate disodium, flavor, microcrystalline cellulose and carboxymethylcellulose sodium, polyethylene glycol, povidone, purified water, sodium phosphate dibasic, sodium phosphate monobasic

Oxymetazoline Hcl 0.05% Nasal Spray
No drip nasal spray

Drug Facts
Active ingredient
Oxymetazoline hydrochloride 0.05%

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Drug Facts (Continued)
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NDC# 68210-5056-1

No Drip Original Pump Mist

Drug Facts (Continued)
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Oxymetazoline Hcl 0.05% Nasal Spray
No drip nasal spray

15 mL

• Original 12 hour pump mist

• fast, powerful congestion relief

MANUFACTURED FOR PRODOSE, INC
2004 Orville Drive North, RONKONKOMA, New York (NY) 11779, United States (USA)

NVZ
LH - 35x26 mm

Batch No :
Mfg. Date :
Expiry Date :
2D Barcode
For Reference Only

OXYMETAZOLINE HYDROCHLORIDE

oxymetazoline hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68210-5056
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.05 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
WATER (UNII: 059QF0KO0R)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SODIUM PHOSPHATE, DIBASIC (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC (UNII: 3980JIH2SW)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	LEMON	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210-5056-1	1 in 1 CARTON	09/03/2025	
1		15 mL in 1 BOTTLE, PUMP; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/03/2025	

Labeler - Prodose, Inc. (119371190)

Revised: 2/2026

Prodose, Inc.