

**OXYMETAZOLINE HYDROCHLORIDE- oxymetazoline hydrochloride spray**  
**Prodose, Inc.**

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**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.

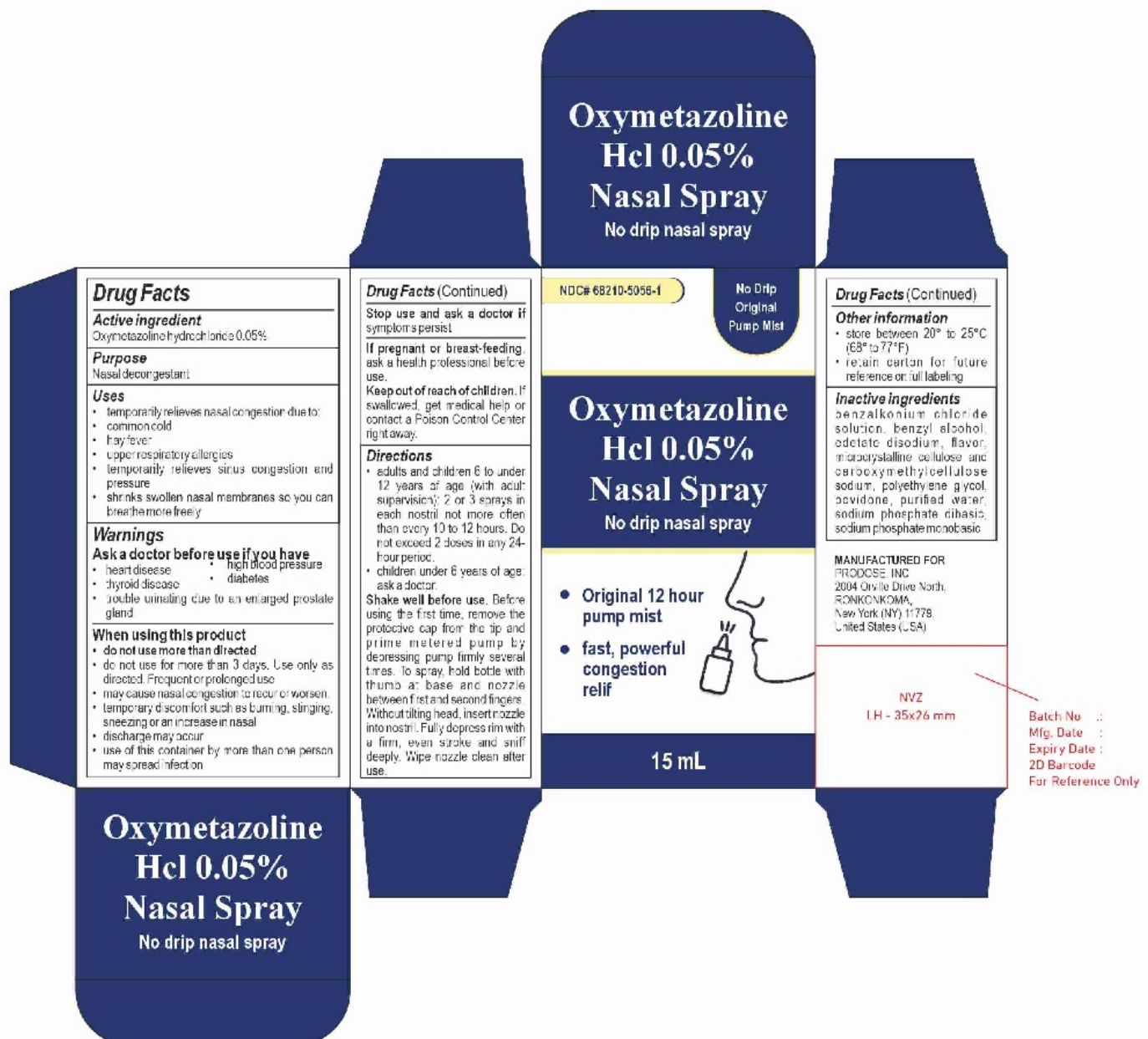
### Other information

- 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

### PRINCIPAL DISPLAY PANEL - 15 mL Bottle Carton



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	
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**Labeler -** Prodose, Inc. (119371190)

Revised: 9/2025

Prodose, Inc.