

**QUALITY CHOICE NASAL MIST SEVERE CONGESTION- oxymetazoline hydrochloride spray**  
**CHAIN DRUG MARKETING ASSOCIATION**

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**Quality Choice Nasal Mist Severe Congestion 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:** Shake well before use. Push down cap while turning counter-clockwise and remove cap. Remove clip under rim. Before using the first time, prime metered pump by depressing pump firmly several times. To spray, hold bottle with thumb at base and nozzle rim 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. Replace clip under rim and secure cap after use.

**Other information**

- store at room temperature

**TAMPER EVIDENT: DO NOT USE IF PRINTED SEAL OVER CAP IS BROKEN OR MISSING**

**How to use:**

Push down cap while turning counter-clockwise and remove cap. Remove clip under rim. Secure cap after use.

**Inactive ingredients**

benzalkonium chloride, benzyl alcohol, camphor, edetate disodium, eucalyptol, glycerin, menthol, microcrystalline cellulose and carboxymethylcellulose sodium, polyethylene glycol, povidone, propylene glycol, purified water, sodium phosphate dibasic, sodium phosphate monobasic, xanthan gum.

**Package/Label Principal Display Panel**

QUALITY CHOICE ®

NDC# **83324-274-01**

**\*Compare to Active Ingredient Afrin® No Drip Severe Congestion Pump Mist**

No Drip

12 Hour Relief

**Nasal Mist**

**Severe Congestion**

Oxymetazoline HCl 0.05%

Nasal Solution

Fast, Powerful Congestion Relief

For Colds & Allergies

No Drip Pump Mist Won't Drip from Nose or Down Throat

**Plus Menthol**

1 FL OZ (30mL)

100% QC SATISFACTION GURANTEED

Distributed by C.D.M.A., Inc.

Novi, MI 48375

[www.qualitychoice.com](http://www.qualitychoice.com)

Questions: 800-935-2362

\*This product is not manufactured or distributed by Bayer Healthcare LLC, distributor of Afrin<sup>®</sup> No Drip Severe Congestion Pump Mist.



## QUALITY CHOICE NASAL MIST SEVERE CONGESTION

oxymetazoline hydrochloride spray

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-274
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)				
BENZYL ALCOHOL (UNII: LKG8494WBH)				
CAMPHOR (NATURAL) (UNII: N20HL7Q941)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
EUCALYPTOL (UNII: RV6J6604TK)				
GLYCERIN (UNII: PDC6A3C0OX)				
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)				
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JIH2SW)				
XANTHAN GUM (UNII: TTV12P4NEE)				
Product Characteristics				
Color	white (off white)	Score		
Shape		Size		
Flavor	MENTHOL	Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-274-01	1 in 1 CARTON	05/02/2025	
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	05/02/2025	

**Labeler** - CHAIN DRUG MARKETING ASSOCIATION (011920774)

