QUALITY CHOICE- oxymetazoline hydrochloride spray
CHAIN DRUG MARKETING ASSOCIATION

Quality Choice 12 Hour Relief Nasal Spray Drug Facts

Active ingredient

Oxymetazoline hydrochloride 0.05%

Purpose

Nasal Decongestant

Uses

temporarily relieves nasal congestion due to:

- a cold
- hay fever
- upper respiratory allergies
- promotes nasal and sinus drainage
- temporarily relieves sinus congestion and pressure
- helps clear nasal passages; shrinks swollen nasal membranes

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 (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

Shake well before use: Before using the first time, remove the protective cap from the tip by pressing and twisting off the cap.

Rotate the lock tab to align arrow marks to unlock the pump. Prime the lock tab to align arrow marks to unlock the pump. Prime metered pump by depressing pump firmly several times.

To spray, hold bottle with thump 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. Lock the pump by ensuring that arrow marks are not aligned and replace the protective cap on tip.

Other information

- store at room temperature
- retain carton for future reference on full labeling

Inactive ingredients

benzalkonium chloride, benzyl alcohol, camphor, edetate disodium, eucalyptol, menthol, polysorbate 80, propylene glycol, purified water, sodium phosphate dibasic, sodium phosphate monobasic.

Questions or comments?

1-866-467-2748

Principal Display Panel

NDC# **63868-699-22**

QUALITY CHOICE ®

* **Compare to Active Ingredient in Mucinex® Full Force Nasal Spray**

12 Hour Relief

Nasal Spray

Nasal Decongestant

Oxymetazoline HCl 0.05%

Fast Relief of Sinus Pressure & Nasal Congestion

3/4 FL OZ (22 mL)

100% QC SATISFACTION GUARANTEED

Distributed by C.D.M.A. Inc. ©

43157 W 9 Mile Rd

Novi, MI 48375

www.qualitychoice.com

Questions: 248-449-9300

*This product is not manufactured or distributed by Reckitt Benckiser, the distributor of Mucinex ®Full Force™ Nasal Spray.

QUALITY CHOICE

oxymetazoline hydrochloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-699
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)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)	
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JIH2SW)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-699-22	1 in 1 CARTON	12/17/2018	
1		22 mL in 1 BOTTLE, PUMP; 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	12/17/2018	

Labeler - CHAIN DRUG MARKETING ASSOCIATION (011920774)

Revised: 12/2025 CHAIN DRUG MARKETING ASSOCIATION