TOPCARE NASAL- oxymetazoline hydrochloride spray Topco Associates LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Topco Associates LLC. Nasal Spray 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 (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 20-25°C (68-77°F)
- retain carton for future reference on full labeling

Inactive ingredients

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

Questions or comments?

1-888-423-0139

Principal Display Panel

COMPARE TO AFRIN® ALLERGY SINUS NASAL SPRAY ACTIVE INGREDIENT

Nasal Spray

NASAL DECONGESTANT OXYMETAZOLINE HCl 0.05%

ALLERGY SINUS

12 HOUR RELIEF

OUR PHARMACISTS RECOMMEND

Fast, Powerful Congestion Relief from Allergies

Reduces Swelling of Nasal Passages

1 FL OZ (30 mL)





COMPARE TO AFRIN® ALLERGY SINUS NASAL SPRAY ACTIVE INGREDIENT

Nasal Spray

NASAL DECONGESTANT OXYMETAZOLINE HCI 0.05%

ALLERGY SINUS





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- Reduces Swelling of Nasal Passages

1 FL OZ (30 mL)





IS BROKEN OR MISSING DO NOT USE IF PRINTED NECKBAND



Nasal

OXYMETAZOLINE HCI 0.05%

ALLERGY SINUS



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This TopCare® product is laboratory tested to guarantee its highest quality. Your total satisfaction is guaranteed.

*This product is not manufactured or distributed by Bayer HealthCare LLC, distributor of Afrin® Allergy Sinus Nasal Spray.

> SUSTAINABLE FORESTRY INITIATIVE Certified Sourcing

Drug Facts

Active ingredient

Purpose Oxymetazoline hydrochloride 0.05%......Nasal decongestant

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Drug Facts (continued)

Inactive ingredients benzalkonium chloride solution. benzyl alcohol, camphor, dibasic sodium phosphate, edetate disodium, eucalyptol, menthol, monobasic sodium phosphate, polysorbate 80, propylene glycol,

Questions or comments? 1-888-423-0139





NDC:36800-817

81710 88 C6

TOPCARE NASAL

oxymetazoline hydrochloride spray

Product Information

HUMAN OTC DRUG Product Type Item Code (Source)

NASAL Route of Administration

Active Ingredient/Active Moiety

Basis of Strength Ingredient Name Strength

OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE -	OXYMETAZOLINE	0.05 g
UNII:8 VLN5B44ZY)	HYDROCHLORIDE	in 100 mL

Inactive Ingredients			
Ingredient Name	Strength		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)			
BENZYL ALCOHOL (UNII: LKG8494WBH)			
SODIUM PHO SPHATE, DIBASIC (UNII: GR686LBA74)			
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)			
EUCALYPTOL (UNII: RV6J6604TK)			
MENTHOL (UNII: L7T10EIP3A)			
SODIUM PHO SPHATE, MO NO BASIC (UNII: 3980 JIH2SW)			
POLYSORBATE 80 (UNII: 6OZP39ZG8H)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
1 NDC:36800-817-10	1 in 1 CARTON	08/06/1996		
1	30 mL in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part341	08/06/1996		

Labeler - Topco Associates LLC (006935977)

Revised: 11/2018 Topco Associates LLC