MAXIMUM STRENTH NASAL- oxymetazoline hydrochloride - 0.05% liquid Walgreens

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.

Walgreens

Drug Facts

Active ingredient

Purpose

Oxymetazoline Hydrochloride - 0.05%

Nazal Decongestant

Uses

For the temporary relief of nasal congestion due to the common cold, hay fever or other upper respiratory allergies.

Warnings

Do not use this product if person has heart disease, high blood pressure, thyroid disease, diabetes, or if adult has difficulty in urination due to enlargement of the prostate gland unless directed by a doctor.

When using this product

Do not exceed recommended dosage

Do not use this product for more than 3 days. Use only as directed. Frequent or prolonged use may cause nasal congestion to recur or worsen. If symptoms persist, consult a doctor. This product may cause temporary discomfort such as burning, stinging, sneezing, or an increase in nasal discharge. The use of this container by more than one person may spread infection.

Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away

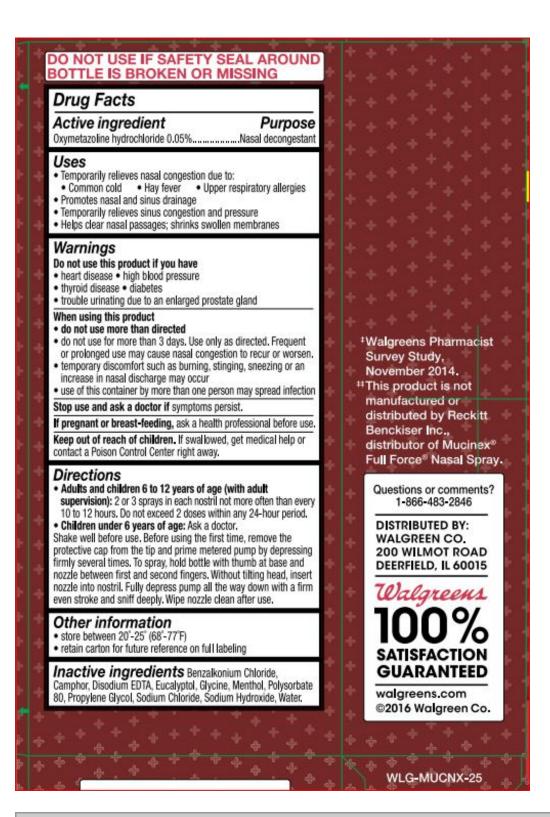
Directions

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**: consult a doctor.

Inactive ingredients

Benzalkonium Chloride Camphor Disodium EDT A Eucalyptol Glycine Menthol Polysorbate 80 Propylene Glycol Sodium Chloride





MAXIMUM STRENTH NASAL

oxymetazoline hydrochloride - 0.05% liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-7003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
	Oxymetazoline Hydrochloride	0.05 g in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
Benzalkonium Chloride (UNII: F5UM2KM3W7)		
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)		
EDETATE DISO DIUM (UNII: 7FLD91C86K)		
Eucalyptol (UNII: RV6J6604TK)		
Glycine (UNII: TE7660 XO1C)		
Menthol (UNII: L7T10EIP3A)		
Polysorbate 80 (UNII: 6OZP39ZG8H)		
Propylene Glycol (UNII: 6DC9Q167V3)		
Sodium Chloride (UNII: 451W47IQ8X)		
Sodium Hydroxide (UNII: 55X04QC32I)		
water (UNII: 059QF0KO0R)		

Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
П	NDC:0363-7003-01	1 in 1 CARTON	07/15/2016	
	1	22 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	07/15/2016	

Labeler - Walgreens (008965063)

Registrant - Product Quest Mfg. (927768135)

Establishment				
Name	Address	ID/FEI	Business Operations	
Product Quest Mfg.		927768135	manufacture(0363-7003), label(0363-7003)	

Revised: 2/2018 Walgreens