MAX STR.NASAL RELIEF SEVERE CONGESTION- oxymetazoline hydrochloride 0.05% liquid
Ride Aid

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.

Drug Facts

Active ingredient	**Purpose**
Oxymetazoline Hydrochloride 0.05%	Nasal decongestant

Uses

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

Warnings

Ask a doctor before use if you have
• heart disease• high blood pressure• thyroid disease
• diabetes• trouble urinating due to enlarged prostate gland

When using this product

Do not use this product if you have heart disease • high blood pressure • thyroid disease • diabetes • or difficulty in urination due to enlargement of the prostate gland
When using this product • Do not exceed recommended dosage. • 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. • Do not use for more than 3 days. Use only as directed. • Frequent or prolonged use may cause nasal congestion to recur or worsen.
Stop use and ask doctor if symptoms persist, consult a doctor.

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

Directions

Before using the first time, remove the protective cap from the tip and prime 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. • adults and children 6 to under12 years of age (with adult supervision): 2 or 3 sprays in each nostril not more often than every 10 to 1 2 hours. Do not exceed 2 doses within any 24-hour period
• children under 6 years of age: consult a doctor

Inactive ingredients

Benzalkonium Chloride, Benzyl Alcohol, Camphor, Edetate Disodium, Eucalyptol, Menthol, Microcrystalline Cellulose, Polyethylene Glycol, Povidone, Propylene Glycol, Purified Water, Sodium Carboxymethyl Cellulose, Sodium Phosphate Dibasic, Sodium Phosphate Monobasic.

RITE AID PHARMACY

nasal relief

severe congestion nasal spray

oxymetazoline hydrochloride 0.05%

RITE AID PHARMACY

*Compare to the active ingredient in Afrin® No-Drip®

MAXIMUM STRENGTH

nasal relief

severe congestion nasal spray

oxymetazoline hydrochloride 0.05%

NASAL DECONGESTANT

12 HOUR

won't drip from nose or down throat

fast, powerful congestion relief

colds; allergies

no drip pump spray

1 FL OZ (30 mL)

CAP REMOVAL INSTRUCTIONS:
To remove cap squeeze cap at base on opposite sides on the two smooth impressions with fingers. While squeezing, twist cap counterclockwise to remove. Refer to directions on top of cap.

To reapply cap turn cap on pump clockwise until it locks.

DISTRIBUTED BY:
RITE AID
30 HUNTER LANE
CAMP HILL, PA 17011

SATISFACTION GUARANTEE
RITE AID
IF YOU'RE NOT SATISFIED, WE'LL HAPPILY REFUND YOUR MONEY.

ACTUAL SIZE

MAX STR.NASAL RELIEF SEVERE CONGESTION

oxymetazoline hydrochloride 0.05% liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11822-1232
Route of Administration	TOPICAL		

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 (SYNTHETIC) (UNII: 5TJD82A1ET)	
Edetate Disodium (UNII: 7FLD91C86K)	
Eucalyptol (UNII: RV6J6604TK)	
Menthol (UNII: L7T10EIP3A)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
Povidone (UNII: FZ989GH94E)	
Propylene Glycol (UNII: 6DC9Q167V3)	
Water (UNII: 059QF0KO0R)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED FORM (UNII: K679OBS311)	
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS (UNII: 22ADO53M6F)	
SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS (UNII: KH7I04HPUU)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-1232-1	1 in 1 CARTON	05/09/2014	
1		15 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	05/09/2014	

Labeler - Ride Aid (014578892)

Registrant - Product Quest Mfg, LLC (927768135)

Establishment

Name	Address	ID/FEI	Business Operations
Product Quest Mfg, LLC		927768135	manufacture(11822-1232) , label(11822-1232)