

NASAL- oxymetazoline hydrochloride spray
Rite Aid Corporation

Rite Aid Corporation Nasal Spray Drug Facts

Active ingredient

Oxymetazoline HCl 0.05%

Purpose

Nasal decongestant

Uses

temporarily relieves

- nasal congestion due to a cold, hay fever, or other upper respiratory allergies
- sinus congestion and pressure

Warnings

Ask a doctor before use if you have

- heart disease
- thyroid disease
- diabetes
- high blood pressure
- trouble urinating due to an enlarged prostate gland

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.
- 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

Shake well before use. Hold white tabs, SQUEEZE grooved area of cap FIRMLY and turn counter clockwise. Before using for the first time, prime the pump by firmly depressing its rim several times. Hold container with thumb at base and nozzle between first and second fingers. Without tilting your head, insert nozzle into nostril. Fully depress rim with a firm, even stroke and inhale deeply. Secure cap after use.

adults & children 6 yrs. & older (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 24 hours.
children 2 to under 6 yrs.	ask a doctor
children under 2 yrs.	do not use

Other information

- store at 20-25°C (68-77°F)

Inactive ingredients

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

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

WITH MENTHOL

Compare to the active ingredient of Vicks[®] Sinex[®] Severe

MAXIMUM STRENGTH NASAL SPRAY

NASAL DECONGESTANT

OXYMETAZOLINE HCl

SEVERE

Fast & powerful relief

Sinus congestion & pressure

ULTRA FINE MIST

12 HOUR

1 FL OZ (30 mL)

DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING

DISTRIBUTED BY:
RITE AID
200 NEUBERRY COMMONS
ETTERS, PA 17319
www.riteaid.com

SATISFACTION GUARANTEE
If you're not satisfied, we'll happily refund your money.

Only selected information is listed on the bottle label. Keep this carton for future reference.

WITH MENTHOL NDC 11822-6623-1

Compare to the active ingredient of
Vicks® Sinex® Severe*

MAXIMUM STRENGTH
NASAL SPRAY

NASAL DECONGESTANT

OXYMETAZOLINE HCl
SEVERE

Fast & powerful relief
Sinus congestion & pressure

ULTRA FINE MIST

12 HOUR

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ACTUAL SIZE

CODE AREA
7X010 83 C2

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*This product is not manufactured or distributed by Procter & Gamble, distributor of Vicks® Sinex® Severe.

How to use:
Hold white tabs, **SQUEEZE** grooved area of cap **FIRMLY** and turn counter clockwise. Secure cap after use.

SQUEEZE grooves **FIRMLY** while turning

Hold white tabs **FIRMLY**

NASAL

oxymetazoline hydrochloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-6623
Route of Administration	NASAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	.05 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
EUCALYPTOL (UNII: RV6J6604TK)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM (UNII: 3980JIH2SW)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	WHITE (off white)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-6623-1	1 in 1 CARTON	12/05/2018	
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	12/05/2018	

Labeler - Rite Aid Corporation (014578892)

Revised: 7/2024

Rite Aid Corporation