SINUFRIN QUICK RELIEF DECONGESTANT- sinufrin spray
Neilmed pharmaceuticals Inc.

NeilMed Sinufrin Quick Relief Decongestant

Drug Facts
Active Ingredients

Oxymetazoline hydrochloride 0.04%

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
- Diabetes
- Thyroid disease
- 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.

Warnings

Stop use and ask a doctor if symptoms persist

Warnings

If pregnant or breast-feeding, ask a health professional before use.

Warnings

Keep out of reach of children. If 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): 3 or 4 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 between 68° F (20° C) and 86° F (30° C)
- Retain carton for future reference on full labeling

Inactive Ingredients

Sodium chloride, sodium bicarbonate, propylene glycol, edetate disodium, benzalkonium chloride, purified water.

Principal Display Panel

SINUFRIN QUICK RELIEF DECONGESTANT

sinufrin spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:13709-325
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OXYMETAZOLINE (UNII: 8VLN5B44ZY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE	0.4 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:13709-325-01	1 in 1 CARTON	12/11/2023	
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 Drug	M012	12/11/2023	

Labeler - Neilmed pharmaceuticals Inc. (799295915)

Establishment

Name	Address	ID/FEI	Business Operations
NeilMed Pharmaceuticals Inc.		799295915	manufacture(13709-325)

Revised: 1/2024 Neilmed pharmaceuticals Inc.