VICKS SINEX SEVERE ORIGINAL- oxymetazoline hydrochloride spray
Procter & Gamble Manufacturing GmbH

Vicks ® Sinex™

Severe Original 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 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 increased 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

Directions

adults & children 6 years & 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 years	ask a doctor
children under 2 years	do not use

Other information

- do not exceed 25ºC

Inactive ingredients

benzalkonium chloride, benzyl alcohol, citric acid anhydrous, edetate disodium, fragrance, polysorbate 80, propylene glycol, purified water, sodium citrate

Questions?

1-800-873-8276

TAMPER EVIDENT: Carton sealed for your protection.

MADE IN GERMANY

DIST. BY PROCTER & GAMBLE,

CINCINNATI OH 45202

PRINCIPAL DISPLAY PANEL - 15 ml Bottle Carton

VICKS ®

Sinex™

SEVERE

Oxymetazoline HCl Nasal Decongestant

ORIGNAL

NASAL SPRAY

- Fast Sinus Congestion & Pressure Relief
- Powerful Vicks Vapors

12
HOUR

½ FL OZ
(15 ml)

VICKS SINEX SEVERE ORIGINAL

oxymetazoline hydrochloride spray

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:64336-170
Route of Administration	NASAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.0005095 g in 1 mL

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	

Product Characteristics

Color	white (Clear)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:64336-170-01	1 in 1 CARTON	12/20/2018	
1		15 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	12/20/2018	

Labeler - Procter & Gamble Manufacturing GmbH (333608813)

Revised: 10/2024 Procter & Gamble Manufacturing GmbH