VICKS SINEX SCENT FREE- oxymetazoline hydrochloride spray The Procter & Gamble Manufacturing Company

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.

Vicks ® SinexTM

Scent Free

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

Remove protective cap. 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.

adults & children 6	years &	2 or 3 sprays in each nostril, not more
older (with adult		often than every 10 to 12 hours. Do not
supervision)		exceed 2 doses in 24 hours.
children 2 to under	6 years	ask a doctor
children under 2 ye	ars	do not use

Other information

• store at room temperature

Inactive ingredients

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

Questions or comments?

1-800-873-8276

Dist. by Procter & Gamble, Cincinnati OH 45202

PRINCIPAL DISPLAY PANEL - 15 ml Bottle Carton

VICKS ®

 $Sinex^{TM}$ $SCENT\ FREE$

Oxymetazoline HCl - Nasal Decongestant

- Sinus Congestion & Pressure
 - Fast & Powerful Relief

Ultra

Fine

Mist

12

HR

1/2 FL OZ (15 ml)







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

P& $oldsymbol{G}$ www.vicks.com

Made in Czech Republic Dist. by Procter & Gamble, Cincinnati OH 45202

Patents: www.pg.com/patents 90724646 / 1181100







TAMPER EVIDENT: Carton sealed for your protection.

Drug Facts

Active ingredient Oxymetazoline HCI 0.05% Purpose

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 • thyroid disease • diabetes

 heart disease high blood pressure

trouble urinating due to enlarged prostate gland

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adults & children children 2 to under 6 years ask a doctor

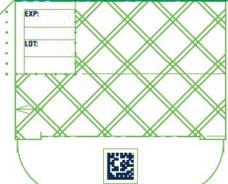
2 or 3 sprays in each nostril, not 6 years & older more often than every 10 to 12 hours (with adult supervision) Do not exceed 2 doses in 24 hours.

children under 2 years do not use

Other information • store at room temperature

Inactive ingredients benzalkonium chloride, benzyl alcohol, citric acid anhydrous, edetate disodium polysorbate 80, purified water, sodium citrate, sorbitol

Questions or comments? 1-800-873-8276





oxymetazoline hydrochloride spray

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37000-435	
Route of Administration	NASAL			

	Active Ingredient/Active Moiety		
ı	Ingredient Name	Basis of Strength	Strength
	OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.05 g in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)		
BENZYL ALCOHOL (UNII: LKG8494WBH)		
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)		
EDETATE DISO DIUM (UNII: 7FLD9 1C86K)		
POLYSORBATE 80 (UNII: 6OZP39ZG8H)		
WATER (UNII: 059QF0KO0R)		
SODIUM CITRATE (UNII: 1Q73Q2JULR)		
SORBITOL (UNII: 506T60A25R)		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37000-435- 01	1 in 1 CARTON	0 1/3 1/2 0 15	
1		15 mL in 1 BOTTLE, PUMP; 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	0 1/3 1/2 0 15	12/01/2021

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 12/2019 The Procter & Gamble Manufacturing Company