

**ULTRA FINE MIST SINUS RELIEF- oxymetazoline hydrochloride spray**  
**DOLGENCORP, INC.**

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**Dollar General Severe Nasal Spray Drug Facts**

**Active ingredient**

Oxymetazoline hydrochloride 0.05%

**Purpose**

Nasal Decongestant

**Uses**

temporarily relieves

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

**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 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 at (1-800-222-1222).

## **Directions**

- **adults & children 6 to under 12 years of age & 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 any 24-hour period.
- **children 2 to under 6 years:** ask a doctor
- **children 2 years:** do not use

**To Use:** Shake well before use. Push down cap while turning counter-clockwise and remove cap. Remove clip under rim. Before using the first time, prime metered 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. Replace clip under rim and secure cap after use.

## **Other information**

- store at room temperature

## **Inactive ingredients**

benzalkonium chloride, benzyl alcohol, camphor, edetate disodium, eucalyptol, menthol, polysorbate 80, propylene glycol, purified water, sodium phosphate dibasic, sodium phosphate monobasic.

## **Questions or comments?**

**1-866-467-2748**

## **Principal Display Panel**

**NDC# 55910-627-30**

**Compare to the active ingredient in Vicks® Sinex™ Severe Original\***

ULTRA FINE MIST

**SINUS RELIEF**

**Oxymetazoline HCl**

Nasal Decongestant

**Original**

- **Fast Sinus Congestion & Pressure Relief**

**12 Hour**

## Pump Mist

1 FL OZ (30 mL)

**IMPORTANT:** Keep the carton for future reference on full labeling

### Distributed by:

\*This product is not manufactured or distributed by Procter and Gamble, the distributor of Vicks <sup>®</sup> Sinex <sup>™</sup> Severe Original



# ULTRA FINE MIST SINUS RELIEF

oxymetazoline hydrochloride spray

## Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-627
Route of Administration	NASAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>OXYMETAZOLINE HYDROCHLORIDE</b> (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.05 g in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)	
<b>CAMPHOR (NATURAL)</b> (UNII: N20HL7Q941)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>EUCALYPTOL</b> (UNII: RV6J6604TK)	
<b>MENTHOL, UNSPECIFIED FORM</b> (UNII: L7T10EIP3A)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM</b> (UNII: GR686LBA74)	
<b>SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM</b> (UNII: 3980JIH2SW)	

## 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:55910-627-30	1 in 1 CARTON	09/18/2020	
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
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OTC Monograph Drug	M012	09/18/2020	
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**Labeler -** DOLGENCORP, INC. (068331990)

Revised: 8/2024

DOLGENCORP, INC.