GOOD SENSE NASAL- oxymetazoline hydrochloride spray L. Perrigo 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.

Perrigo Nasal Spray Drug Facts

Active ingredient

Oxymetazoline hydrochloride 0.05%

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

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

- adults and children 6 to under 12 years of age (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 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 at 20-25°C (68-77°F)
- retain carton for future reference on full labeling

Inactive ingredients

benzalkonium chloride solution, benzyl alcohol, dibasic sodium phosphate, edetate disodium, monobasic sodium phosphate, polyethylene glycol, povidone, propylene glycol, purified water

Questions or comments?

1-800-719-9260

Principal Display Panel

12 Hour Relief

Maximum Strength

Nasal Spray

Oxymetazoline HCl 0.05%

Nasal Decongestant

Original

#1 DOCTOR RECOMMENDED adult nasal spray active ingredient

Fast, Powerful Congestion Relief

For Colds and Allergies

Compare to active ingredient of Afrin® Original Nasal Spray

100% SATISFACTION GUARANTEED

1 FL OZ (30 mL)



12 Hour Relief

Maximum Strength

Oxymetazoline HCI 0.05%

Nasal Decongestant

 Fast, Powerful Congestion Relief

 For Colds and Allergies

1 FL 0Z (30 mL)

Compare to active ingredient of Afrin® Original Nasal Spray



GOODSENSE. 12 Hour Relief

Maximum Strength



Oxymetazoline HC10.05% NasalDecongestant

Gluten Free

Distributed By

Allegan, MI 49010

Drug Facts

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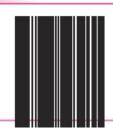
Questions or comments? 1-800-719-9260

Of U.S. physicians surveyed by an independent market research firm.





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GOOD SENSE NASAL

oxymetazoline hydrochloride spray

Pro	duct	Information
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:0113-0304

Route of Administration NASAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

0.5 mg in 1 mL

Inactive Ingredients			
Ingredient Name	Strength		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7)			
BENZYL ALCOHOL (UNII: LKG8494WBH)			
SO DIUM PHO SPHATE, DIBASIC, UNSPECIFIED FORM (UNII: GR686LBA74)			
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)			
SO DIUM PHO SPHATE, MO NO BASIC, UNSPECIFIED FORM (UNII: 3980 JIH2SW)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
PO VIDO NE, UNSPECIFIED (UNII: FZ989GH94E)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			

Product Characteristics					
Color	WHITE (Translucent)	Score			
Shape		Size			
Flavor		Imprint Code			
Contains					

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:0113-0304- 10	1 in 1 CARTON	09/15/1989		
1		30 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product			
2	NDC:0113-0304- 72	1 in 1 CARTON	09/15/1989		
2		37 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 final	part341	09/15/1989				

Labeler - L. Perrigo Company (006013346)

L. Perrigo Company Revised: 12/2019