

## **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.*

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### **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)



GOOD SENSE NASAL			
oxymetazoline hydrochloride spray			
Product Information			
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	

<b>OXYMETAZOLINE HYDROCHLORIDE</b> (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZOLINE HYDROCHLORIDE	0.5 mg in 1 mL
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### Inactive Ingredients

Ingredient Name	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7)	
<b>BENZYL ALCOHOL</b> (UNII: LKG8494WBH)	
<b>SODIUM PHOSPHATE, DIBASIC, UNSPECIFIED FORM</b> (UNII: GR686LBA74)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>SODIUM PHOSPHATE, MONOBASIC, UNSPECIFIED FORM</b> (UNII: 3980JIH2SW)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Product Characteristics

<b>Color</b>	WHITE (Translucent)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### 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)