# GOOD SENSE NASAL DECONGESTANT- pseudoephedrine hcl tablet, film coated 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 Decongestant Drug Facts**

### **Active ingredient (in each tablet)**

Pseudoephedrine HCl 30 mg

### Purpose

Nasal decongestant

### Uses

- temporarily relieves sinus congestion and pressure
- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies

### Warnings

### Do not use

if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

### Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland

### When using this product

do not exceed recommended dosage

### Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- symptoms do not improve within 7 days or occur with fever

### If pregnant or breast-feeding,

ask a health professional before use.

### Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

### **Directions**

adults and children 12 years and over	<ul> <li>take 2 tablets every 4 to 6 hours</li> <li>do not take more than 8 tablets in 24 hours</li> </ul>
children ages 6 to 11 years	<ul> <li>take 1 tablet every 4 to 6 hours</li> <li>do not take more than 4 tablets in 24 hours</li> </ul>
children under 6 years	do not use this product in children under 6 years of age

### Other information

- each tablet contains: calcium 20 mg
- store at 20°-25°C (68°-77°F)
- do not use if blister unit is broken or torn

### **Inactive ingredients**

carnauba wax, dibasic calcium phosphate dihydrate, FD&C red no. 40 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, silicon dioxide, titanium dioxide

### Questions or comments?

1-800-719-9260

### **Principal Display Panel**

Maximum Strength

Congestion

Non-Drowsy

Nasal Decongestant

Actual Size

Pseudoephedrine HCl Tablets

Nasal Decongestant

Nasal & Sinus Congestion

Sinus Pressure

Compare to active ingredient of Sudafed® Congestion Tablets

24 Tablets (30 mg Each)

# GOODSENSE.

# GOODSENSE.

NDC 0113-0432-62

**Maximum Strength** 

# Congestion

Pseudoephedrine HCl Tablets Nasal Decongestant

Actual Size

- Nasal & Sinus Congestion
- Sinus Pressure

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Important: Rea dialiproduct information before using . Keep this box for important information .

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### Drug Facts (continued)

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### Gluten Free

Distributed By



Allegan, MI 49010







### GOOD SENSE NASAL DECONGESTANT

pseudoephedrine hcl tablet, film coated

### **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0113-0432
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Route of Administration ORAL

# Active Ingredient/Active Moiety Ingredient Name Basis of Strength PSEUDO EPHEDRINE HYDRO CHLO RIDE (UNII: 6 V9 V2RYJ8 N) (PSEUDO EPHEDRINE HYDRO CHLO RIDE UNII: 7 CUC 9 DD 19 F) PSEUDO EPHEDRINE HYDRO CHLO RIDE 30 mg

Inactive Ingredients		
Ingredient Name	Strength	
CARNAUBA WAX (UNII: R12CBM0EIZ)		
DIBASIC CALCIUM PHO SPHATE DIHYDRATE (UNII: O7TSZ97GEP)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POLYSORBATE 80 (UNII: 6OZP39ZG8H)		
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		

Product Characteristics			
Color	RED	Score	no score
Shape	ROUND (convex)	Size	7mm
Flavor		Imprint Code	L432
Contains			

Packaging			
# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
NDC:0113-0432-62	1 in 1 CARTON	09/15/1989	
1	24 in 1 BLISTER PACK; 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)

Revised: 12/2019 L. Perrigo Company