NASAL DECONGESTANT- pseudoephedrine hydrochloride tablet, film coated, extended release
Meijer Distribution Inc

Meijer Distribution, Inc. Nasal Decongestant Drug Facts

Active ingredient (in each tablet)

Pseudoephedrine HCl 120 mg

Purpose

Nasal decongestant

Uses

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

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 a 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	• take 1 tablet every 12 hours • do not take more than 2 tablets in 24 hours
children under 12 years	do not use this product in children under 12 years of age

Other information

- **each tablet contains:** calcium 45 mg
- store at 20-25°C (68-77°F) in a dry place
- protect from light
- **do not use if blister unit is broken or torn**
- see carton end panel for lot number and expiration date
- this product meets the requirements of USP Drug Release Test 3

Inactive ingredients

carnauba wax, colloidal silicon dioxide, dibasic calcium phosphate dihydrate, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, titanium dioxide

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

Compare to Sudafed® Sinus Congestion 12 Hour active ingredient

NON-DROWSY

MAXIMUM STRENGTH

nasal decongestant

Pseudoephedrine Hydrochloride Extended-Release Tablets, 120 mg

Nasal Decongestant

12 HOUR SINUS

10 Coated Caplets**

**capsule-shaped tablets

actual size

Sinus Pressure

Congestion

nasal decongestant

Pseudoephedrine Hydrochloride Extended-Release Tablets, 120 mg

Nasal Decongestant

meijer

NDC 41250-809-52

Compare to Sudafed® Sinus Congestion 12 Hour active ingredient*

NON-DROWSY

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nasal decongestant

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Nasal Decongestant

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10 Coated Caplets**

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Sinus Pressure | Congestion

05452 6E C11

COD01-74NR-74

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Drug Facts (continued)

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*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Sudafed®.

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GRAND RAPIDS, MI 49544
www.meijer.com

GF GLUTEN FREE

OUR QUALITY GUARANTEE

WWW.MEIJER.COM/SATISFACTION

PID 533069

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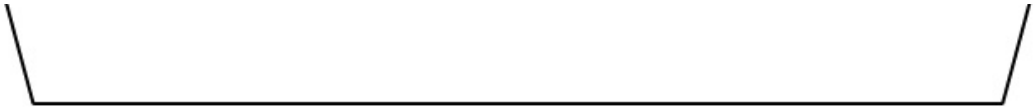

NASAL DECONGESTANT

pseudoephedrine hydrochloride tablet, film coated, extended release

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41250-809
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	120 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CARNAUBA WAX (UNII: R12CBM0EIZ)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	WHITE	**Score**	no score
Shape	OVAL	**Size**	18mm
Flavor		**Imprint Code**	L054
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:41250-809-52	10 in 1 CARTON	11/28/2018	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:41250-809-60	20 in 1 CARTON	11/28/2018	
2		1 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
ANDA	ANDA075153	11/28/2018	

Labeler - Meijer Distribution Inc (006959555)

Revised: 5/2019 Meijer Distribution Inc