GOOD SENSE NASAL DECONGESTANT PE- phenylephrine hydrochloride tablet 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 Decongestant PE Drug Facts

Active ingredient (in each tablet)

Phenylephrine hydrochloride 10 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 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 4 hoursdo not take more than 6 tablets in 24 hours
children under 12 years	ask a doctor

Other information

- store at 20-25°C (68-77°F)
- do not use if blister unit is broken or torn

Inactive ingredients

anhydrous dibasic calcium phosphate, carnauba wax, FD&C red no. 40 aluminum lake, lecithin, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, silicon dioxide, talc, titanium dioxide

Questions or comments?

1-800-719-9260

Principal Display Panel

Maximum Strength

Congestion

Non-Drowsy

Nasal Decongestant PE

Actual Size

1 Pill/Dose

Phenylephrine HCl Tablets

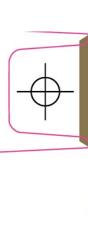
Nasal Decongestant

Nasal & Sinus Congestion

Sinus Pressure

Compare to active ingredient of Sudafed PE® Congestion Tablets

72 Tablets (10 mg Each)



GOODSENSE.

Congestion Non-Drowsy

Nasal Decongestant PE

GOODSENSE_®

NDC 0113-0094-23

Maximum Strength

Congestion Non-Drowsy

Nasal Decongestant PE

Phenylephrine HCl Tablets Nasal Decongestant



- Nasal & Sinus Congestion
- Sinus Pressure

72 Tablets (10 mg Each)

Compare to active ingredient of Sudafed PE® Congestion Tablets





OPEN OTHER END

Important: Read all product information before using. Keep this box for important information.

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Drug Facts (continued)

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Questions or comments? 1-800-719-9260

Gluten Free

Distributed By



Allegan, MI 49010







GOOD SENSE NASAL DECONGESTANT PE

phenylephrine hydrochloride tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0113-0094

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

PHENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)

PHENYLEPHRINE HYDROCHLORIDE

10 mg

Inactive Ingredients

Ingredient Name Strength
ANHYDROUS DIBASIC CALCIUM PHO SPHATE (UNII: L11K75P92J)

CARNAUBA WAX (UNII: R12CBM0EIZ)

MAGNESIUM STEARATE (UNII: 70097M6I30)

MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)

POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)

SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)

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TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Product Characteristics					
Color	RED	Score	no score		
Shape	ROUND	Size	8 mm		
Flavor		Imprint Code	L7		
Contains					

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:0113-0094-23	72 in 1 CARTON	06/28/2005			
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product				
2	NDC:0113-0094-68	36 in 1 CARTON	08/01/2005			
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product				
3	NDC:0113-0094-89	18 in 1 CARTON	06/21/2005			
3		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	
OTC monograph final	part341	06/21/2005		

Labeler - L. Perrigo Company (006013346)

Revised: 12/2019 L. Perrigo Company