HEALTH MART NASAL DECONGESTANT PE- phenylephrine hydrochloride tablet, film coated

Strategic Sourcing Services LLC

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.

McKesson Nasal Decongestant PE Drug Facts

Active ingredient (in each tablet)

Phenylephrine HCl 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 hours do 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

Compare to Sudafed $PE^{\mathbb{R}}$ Congestion

active ingredient

MAXIMUM STRENGTH

Nasal Decongestant PE

Phenylephrine HCl Tablets, 10mg

Nasal & sinus congestion - sinus pressure

ACTUAL SIZE

Non-Drowsy

Gluten Free

18 tablets 10mg each



NDC 62011-0077-2

Compare to Sudafed PE® Congestion active ingredient*

MAXIMUM STRENGTH

Nasal Decongestant PE

Phenylephrine HCI Tablets, 10mg

Nasal & sinus congestion • sinus pressure

ACTUAL SIZE

Non-Drowsy **Gluten Free**

18 tablets 10mg each

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

Drug Facts

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Drug Facts (continued)

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Questions or comments? 1-800-719-9260

*This product is not manu factured or distributed by McNeil Consumer Heal thcare, distributor of Sudafed PE® Congestion.

M⊆KESSON

Another Quality Product Distributed by McKesson 6555 State Highway 161, Irving, TX 75039 Morey Back Guarantee healthmart.com







HEALTH MART NASAL DECONGESTANT PE

phenylephrine hydrochloride tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62011-0077
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	
CARNAUBA WAX (UNII: R12CBM0 EIZ)		
ANHYDRO US DIBASIC CALCIUM PHO SPHATE (UNII: L11K75P92J)		
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)		
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				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62011-0077-1	36 in 1 CARTON	12/23/2011	
1		$1\ \text{in}\ 1\ \text{BLISTER}$ PACK; Type 0: Not a Combination Product		
2	NDC:62011-0077-2	18 in 1 CARTON	08/13/2013	
2		$1\ \text{in}\ 1\ \text{BLISTER}$ 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	12/23/2011	

Labeler - Strategic Sourcing Services LLC (116956644)

Revised: 12/2019 Strategic Sourcing Services LLC