NASAL DECONGESTANT PE- phenylephrine hydrochloride tablet, film coated H E B

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.

HEB 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 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

Compare to Sudafed PE® Congestion active ingredient

Maximum Strength

Nasal Decongestant PE

Phenylephrine HCl Tablets / Nasal Decongestant

Nasal & Sinus Congestion

Non-Drowsy

Relief of:

Nasal & Sinus Congestion

Sinus Pressure

actual size

18 TABLETS, 10 mg EACH

Compare to Sudafed PE® Congestion active ingredient*

NDC 37808-209-89



Maximum Strength

Nasal Decongestant PE

Phenylephrine HCI Tablets / Nasal Decongestant

Nasal & Sinus Congestion

Non-Drowsy

Relief of:

- Nasal & Sinus Congestion
- Sinus Pressure

18 TABLETS, 10 mg EACH

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

Drug Facts

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Drug Facts (continued)

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*This product is not manufactured or distributed by McNeil Consumer Health care, distributor of Sudafed PE® Congestion.

GLUTEN FREE

CONVENIENT RECLOSING TAB











NASAL DECONGESTANT PE

phenylephrine hydrochloride tablet, film coated

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-209

Route of Administration ORAL

Active Ingredient/Active Moiety

3		
Ingredient Name	Basis of Strength	Strength
	PHENYLEPHRINE HYDROCHLORIDE	10 mg

Ingredient Name

CARNAUBA WAX (UNII: R12CBM0EIZ)

ANHYDRO US DIBASIC CALCIUM PHOSPHATE (UNII: L11K75P92J)

MAGNESIUM STEARATE (UNII: 70097M6I30)

MICRO CRYSTALLINE CELLULO SE (UNII: OP1R32D61U)

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)

SILICON DIO XIDE (UNII: ETJ7Z6XBU4)

POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)

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				
ı	# Item Code	:	Package Description	Marketing Star	t Date Marketing End Date
ı	1 NDC:37808-209	0-89 3 in 1 CA	ARTON	09/26/2016	
ı	1	6 in 1 B	LISTER PACK; Type 0: Not a Combination P	Product	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	09/26/2016	

Labeler - HEB (007924756)

Revised: 12/2019 HE B