

SINUS CONGESTION PE- phenylephrine hydrochloride tablet, coated HEB

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.

1131-HEB-2023-0414

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 of age and over: take 1 tablet every 4 hours; do not take more than 6 tablets in 24 hours
- children under 12 years of age: ask a doctor

Other information

- store between 20-25°C (68-77°F) in a dry place
- retain carton for complete product information and warnings

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, D&C red #27 aluminum lake, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, hypromellose, lactose anhydrous, magnesium stearate, polyethylene glycol, stearic acid, titanium dioxide

Questions or comments?

1-844-705-4384

PRINCIPAL DISPLAY PANEL

Compare to Sudafed PE® Sinus Congestion active ingredient†

HEB®

NDC 37808-531-05

Maximum Strength

Sinus Congestion PE

Phenylephrine HCl 10 mg / Nasal Decongestant

Non-Drowsy

Relief of:

- Sinus Pressure • Sinus Congestion

actual size

150 TABLETS - 10 MG EACH

INK AND COATING FREE
FOR LOT AND
EXPIRATION STAMPING



F113105HEB_R0 28960-2003

DO NOT USE IF IMPRINTED SEAL
UNDER CAP IS BROKEN OR MISSING

This product is not manufactured or distributed by Medical Consumer Healthcare, distributor of Sudafed PE® Sinus Congestion.

Drug Facts

Active ingredient (in each tablet) **Purpose**
Phenylephrine HCl 10 mg.....Nasal decongestant

Uses
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temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies

Warnings
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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
nauseousness, dizziness, or sleeplessness occur
symptoms do not improve within 7 days
or occur with a fever

Drug Facts (continued)

If pregnant or breast-feeding, ask a health professional before use.
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Questions or comments?
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DISTRIBUTED BY H-E-B®
SAN ANTONIO, TX 78204

150 TABLETS
10 MG EACH

Maximum Strength
Sinus Congestion PE
Phenylephrine HCl 10 mg
Nasal Decongestant

Compare to Sudafed PE®
Sinus Congestion active ingredient*

NDC 37808-531-05

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• Sinus Pressure
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Sinus Congestion PE
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Non-Drowsy

Relief of:
• Sinus Pressure • Sinus Congestion

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actual size

COATING FREE AREA

SINUS CONGESTION PE

phenylephrine hydrochloride tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-531
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	red	Score	no score
Shape	ROUND	Size	7mm
Flavor		Imprint Code	A;131
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-531-05	1 in 1 CARTON	04/14/2023	
1		150 in 1 BOTTLE, PLASTIC; 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	04/14/2023	

Labeler - HEB (007924756)