

NASAL DECONGESTANT PE- phenylephrine hydrochloride tablet, coated
VALU MERCHANDISERS COMPANY

1131-BST-2021-1117

Drug Facts

Active ingredient (in each tablet)

Phenylephrine HCl 10 mg

Purpose

Nasal decongestant

Uses

- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies, and nasal congestion associated with sinusitis
- 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 use more than directed**

Stop use and ask a doctor if

- you get nervous, dizzy, or sleepless
- symptoms do not improve within 7 days or are accompanied by 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.

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 at 15°-25°C (59°-77°F) in a dry place
- retain carton for complete product information

Inactive ingredients

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

PRINCIPAL DISPLAY PANEL

COMPARE TO THE ACTIVE INGREDIENT IN SUDAFED PE® CONGESTION†

Best Choice®

Non-Drowsy

MAXIMUM STRENGTH

NASAL DECONGESTANT PE

Actual Size

Phenylephrine HCl

For Relief of:

- Sinus Pressure
- Congestion

18 TABLETS

10 mg EACH

NC

NC

NC

Drug Facts (continued)

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Non-Drowsy
MAXIMUM STRENGTH
NASAL DECONGESTANT PE

Best Choice®

COMPARE TO THE ACTIVE
 INGREDIENT IN SUDAFED PE® CONGESTION†

Non-Drowsy
MAXIMUM STRENGTH
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Best Choice®



Phenylephrine HCl
 For Relief of:
 • Sinus Pressure
 • Congestion

Actual Size

18 TABLETS
 10 mg EACH

Best Choice,
 100% Guaranteed
www.bestchoice.us

PROUDLY DISTRIBUTED BY:
 VNU MEDICALS, CO.
 5000 KANSAS AVE
 KANSAS CITY, MO 64116

Report serious side effects to:
 681 Main Street, Lumberton, NJ 08048

**DO NOT USE IF BLISTER UNITS
 ARE TORN OR BROKEN**

Non-Drowsy
MAXIMUM STRENGTH
NASAL DECONGESTANT PE

Best Choice®

NASAL DECONGESTANT PE

phenylephrine hydrochloride tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63941-331
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)	
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:63941-331-02	1 in 1 PACKAGE	03/15/2017	02/28/2025
1		18 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:63941-331-03	2 in 1 CARTON	03/07/2017	02/28/2025
2		18 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 Drug	M012	03/07/2017	02/28/2025

Labeler - VALU MERCHANDISERS COMPANY (868703513)

Revised: 12/2024

VALU MERCHANDISERS COMPANY