

DECONGESTANT PE NON DROWSY- phenylephrine hcl tablet
Allegiant Health

271 - Phenylephrine HCL 5mg

Active ingredient(s)

Phenylephrine HCl 5mg

Purpose

Nasal decongestant

Use(s)

- temporarily relieves sinus congestion and pressure
- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies

Warnings

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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland

When using this product

do not exceed recommended dose

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- symptoms do not improve within 7 days or occur with a fever

Pregnancy/Breastfeeding

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 (1-800-222-1222) right away.

Directions

Adults and children 12 years and over: 1 tablet every 4 hours

- do not take more than 6 tablets in 24 hours

Children under 12 years ask a doctor

Other information

- do not use if imprinted safety seal under cap is broken or missing

Storage

- store at room temperature 20°– 25°C (68°– 77°F)

Inactive ingredients

croscarmellose sodium, lactose, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake,

hypromellose, magnesium stearate, microcrystalline cellulose, silicon dioxide, titanium dioxide.

May contain propylene glycol

Questions

Call 1-888-952-0050 Monday through Friday

Principal Display Panel

HealthA2Z®

NDC 69168-271-17

Non-Drowsy Decongestant PE

Phenylephrine HCl 5 mg • Nasal Decongestant

- Sinus Pressure
- Nasal & Sinus Congestion Due To Colds & Allergies



300 Tablets

Drug Facts

Active ingredient (in each tablet) Purpose
Phenylephrine HCl 5mg.....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

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Drug Facts (continued on inside)



Manufactured by:
Alliant Health
Deer Park, NY 11729

LB1606
R0521B

Questions or comments?
Call 1-888-952-0050 Monday through Friday

Inactive ingredients croscarmellose sodium, lactose, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, hydroxypropyl methylcellulose, magnesium stearate, microcrystalline cellulose, silicon dioxide, titanium dioxide. May contain propylene glycol.

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Adults and children 12 years and over: ■ 1 tablet every 4 hours ■ do not take more than 6 tablets in 24 hours
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days or occur with a fever
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 Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Decongestant PE 5mg

DECONGESTANT PE NON DROWSY

phenylephrine hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69168-271
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

Product Characteristics

Color	red	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	271
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69168-271-17	300 in 1 BOTTLE; Type 0: Not a Combination Product	06/16/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/23/2014	

Labeler - Allegiant Health (079501930)

Revised: 6/2021

Allegiant Health