

**SINUS PRESSURE AND CONGESTION RELIEF PE MAXIMUM STRENGTH-
phenylephrine hcl tablet, film coated
Rite Aid Corporation**

Rite Aid 44-453

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
- 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
- diabetes
- thyroid disease
- high blood pressure
- difficulty in urination due to enlargement of the 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 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 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

- **TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN**
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- see end flap for expiration date and lot number

Inactive ingredients

croscarmellose sodium, dextrose monohydrate, dibasic calcium phosphate dihydrate, FD&C red #40, lecithin, magnesium stearate, maltodextrin, microcrystalline cellulose, silicon dioxide, sodium carboxymethylcellulose, sodium citrate dihydrate, titanium dioxide

Questions or comments?

1-800-426-9391

Principal Display Panel

NDC 11822-4530-2

Compare to the active ingredient in
Sudafed PE® Sinus Congestion*

**MAXIMUM STRENGTH
SINUS PRESSURE &
CONGESTION**

RELIEF PE

PHENYLEPHRINE HCl 10 mg
NASAL DECONGESTANT

Relieves sinus pressure, sinus and nasal congestion
Pseudoephedrine free

NON-DROWSY

ACTUAL SIZE

72
TABLETS

TAMPER EVIDENT: DO NOT USE IF

**PACKAGE IS OPENED OR IF BLISTER
UNIT IS TORN, BROKEN OR SHOWS
ANY SIGNS OF TAMPERING**

*This product is not manufactured or distributed by Johnson & Johnson Corporation,
owner of the registered trademark
Sudafed PE® Sinus Congestion. 50844 REV0820F45323

DISTRIBUTED BY: RITE AID,
200 NEWBERRY COMMONS, ETTERS, PA 17319
www.riteaid.com

SATISFACTION GUARANTEE

If you're not satisfied, we'll
happily refund your money.

MAXIMUM STRENGTH

SINUS PRESSURE & CONGESTION RELIEF PE

PHENYLEPHRINE HCl 10 mg

NDC 11822-4530-2

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0 11822 04176 8

TAMPER EVIDENT: DO NOT USE IF
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no print / no varnish area
lot no & exp date

MAXIMUM STRENGTH
SINUS PRESSURE &
CONGESTION RELIEF PE
PHENYLEPHRINE HCl 10 mg

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SUDAFED PE® SINUS CONGESTION, 50844 REV0820F45323

B-1702-453-23-R3
REV0820F45323

Drug Facts (continued)	Active ingredient (in each tablet) Phenylephrine HCl 10 mg Nasal decongestant
Purpose Nasal decongestant	Uses ■ 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.
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	Ask a doctor before use if you have ■ heart disease ■ diabetes ■ thyroid disease ■ high blood pressure ■ difficulty in urination due to enlargement of the prostate gland
Other information ■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN ■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F) ■ see end flap for expiration date and lot number	When using this product do not exceed recommended dosage. ■ symptoms do not improve within 7 days or occur with fever ■ nervousness, dizziness, or sleeplessness occur Stop use and ask a doctor if
Drug Facts (continued) 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.	Inactive ingredients croscarmellose sodium, dextrose monohydrate, dibasic calcium phosphate dihydrate, FD&C red #40, lecithin, magnesium stearate, malto-dextrin, microcrystalline cellulose, silicon dioxide, sodium carboxymethylcellulose, sodium citrate dihydrate, titanium dioxide
Questions or comments? 1-800-426-9391	

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

Rite Aid 44-453

SINUS PRESSURE AND CONGESTION RELIEF PE MAXIMUM STRENGTH

phenylephrine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-4530
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
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
DEXTROSE MONOHYDRATE (UNII: LX22YL083G)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	red	Score	no score
Shape	ROUND	Size	7mm
Flavor		Imprint Code	44;453
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-4530-7	2 in 1 CARTON	01/14/2005	
1		18 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11822-4530-2	3 in 1 CARTON	01/14/2005	
2		24 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:11822-4530-4	1 in 1 CARTON	01/14/2005	
3		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	01/14/2005	

Labeler - Rite Aid Corporation (014578892)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(11822-4530) , pack(11822-4530)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(11822-4530)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(11822-4530)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(11822-4530)

Revised: 6/2023

Rite Aid Corporation