

**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 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
- 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 11822-4530-7

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

36
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
Kenvue Inc., owner of the registered trademark
Sudafed PE® Sinus Congestion.
50844 REV0820G45307

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

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no print / no varnish area lot no & exp date

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



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

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

*This product is not manufactured or distributed by Kenue Inc., owner of the registered trademark SudaFed PE® Sinus Congestion. RE0820645307 50844

B-1702-453-07-R3 REV0820645307

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

Drug Facts	Active ingredient (in each tablet) Phenylephrine HCl 10 mg Nasal decongestant
Purpose	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 diabetes high blood pressure thyroid disease difficulty in urination due to enlargement of the prostate gland
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	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, lecitin, magnesium stearate, maltodextrin, microcrystalline cellulose, silicon dioxide, sodium carboxymethylcellulose, sodium citrate dihydrate, titanium dioxide
Questions or comments?	1-800-426-9391

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:11822-4530-4	1 in 1 CARTON	01/14/2005	
1		18 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11822-4530-7	2 in 1 CARTON	01/14/2005	
2		18 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:11822-4530-2	3 in 1 CARTON	01/14/2005	
3		24 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: 5/2025

Rite Aid Corporation