

**SINUS PRESSURE AND CONGESTION RELIEF- diphenhydramine hcl,
phenylephrine hcl
Rite Aid Corporation**

Rite Aid 44-453485

Active ingredient (in each tablet) (Sinus Day)

Phenylephrine HCl 10 mg

Purpose

Nasal decongestant

Active ingredients (in each tablet) (Sinus Night)

Diphenhydramine HCl 25 mg

Phenylephrine HCl 10 mg

Purpose

Antihistamine

Nasal decongestant

Uses

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
 - nasal congestion
 - sneezing **(Nighttime only)**
 - runny nose **(Nighttime only)**
 - itchy, watery eyes **(Nighttime only)**
 - itching of the nose or throat **(Nighttime only)**
- temporarily relieves these symptoms due to the common cold:
 - nasal congestion
 - sneezing **(Nighttime only)**
 - runny nose **(Nighttime only)**
- 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.

- with any other product containing diphenhydramine, even one used on skin
(Nighttime only)

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
- a breathing problem such as emphysema or chronic bronchitis **(Nighttime only)**
- glaucoma **(Nighttime only)**

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers. **(Nighttime only)**

When using this product

- **do not exceed recommended dosage**
- excitability may occur, especially in children **(Nighttime only)**
- marked drowsiness may occur **(Nighttime only)**
- alcohol, sedatives, and tranquilizers may increase drowsiness **(Nighttime only)**
- avoid alcoholic beverages **(Nighttime only)**
- use caution when driving a motor vehicle or operating machinery **(Nighttime only)**

Stop use and ask a doctor if

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

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.

Read each section carefully. Do not take DAYTIME and NIGHTTIME products at the same time.

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: do not use

Other information

- **each tablet contains:** calcium 25 mg **(Nighttime only)**
- **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 (Daytime only)

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

Inactive ingredients (Nighttime only)

croscarmellose sodium, dibasic calcium phosphate dihydrate, FD&C blue #1 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, polydextrose, polyethylene glycol, silicon dioxide, stearic acid, titanium dioxide, triacetin

Questions or comments?

1-800-426-9391

Principal display panel

NDC 11822-9485-9

Compare to the active ingredients in **Sudafed PE® Sinus Congestion Day + Night***

SINUS PRESSURE & CONGESTION RELIEF PE

DAYTIME

PHENYLEPHRINE HCl
NASAL DECONGESTANT
NON-DROWSY

Relieves nasal
congestion

ACTUAL SIZE

NIGHTTIME

DIPHENHYDRAMINE HCl
PHENYLEPHRINE HCl
ANTIHISTAMINE,
NASAL DECONGESTANT

Relieves nasal
congestion,
runny nose

ACTUAL SIZE

12 DAYTIME

TABLETS

8 NIGHTTIME

TABLETS

**TAMPER EVIDENT: DO NOT USE IF PACKAGE IS
OPENED OR IF BLISTER UNIT IS TORN, BROKEN
OR SHOWS ANY SIGNS OF TAMPERING**

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.

*The product is not manufactured or distributed by Johnson
& Johnson Corporation, owner of the registered trademark
SUDAFED PE® SINUS CONGESTION DAY + NIGHT

50844 REV0820D45348509

**Do Not Take Daytime and
Nighttime Products at the
Same Time**

B9811R2

<p>Drug Facts (continued)</p> <p>When using this product:</p> <ul style="list-style-type: none"> do not exceed recommended dosage excitability may occur, especially in children (Nighttime only) marked drowsiness may occur (Nighttime only) alcohol, sedatives, and tranquilizers may increase drowsiness (Nighttime only) avoid alcoholic beverages (Nighttime only) use caution when driving a motor vehicle or operating machinery (Nighttime only) <p>Stop use and ask a doctor if:</p> <ul style="list-style-type: none"> symptoms do not improve within 7 days or occur with fever nervousness, dizziness, or sleeplessness occur <p>If pregnant or breast-feeding, ask a health professional before use.</p> <p>Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.</p> <p>Read each section carefully. Do not take DAYTIME and NIGHTTIME products at the same time.</p>	<p>Drug Facts (continued)</p> <p>Directions</p> <ul style="list-style-type: none"> 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: do not use <p>Other information</p> <ul style="list-style-type: none"> each tablet contains: calcium 25 mg (Nighttime only) TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BUSTER IS TORN OR BROKEN store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F) use end flap for expiration date and lot number <p>Inactive ingredients (Daytime only)</p> <p>croscarmellose sodium, dextrose monohydrate, dibasic calcium phosphate dihydrate, FD&C red #40, lecithin, magnesium stearate, methylcellulose, microcrystalline cellulose, silicon dioxide, sodium carboxymethylcellulose, sodium citrate dihydrate, titanium dioxide</p>
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<p>Drug Facts</p> <p>Active ingredient (in each tablet) (Sinus Day)</p> <p>Phenylephrine HCl 10 mg</p> <p>Purpose</p> <p>Nasal decongestant</p> <p>Active ingredients (in each tablet) (Sinus Night)</p> <p>Diphenhydramine HCl 25 mg</p> <p>Antihistamine</p> <p>Phenylephrine HCl 10 mg</p> <p>Nasal decongestant</p> <p>Uses</p> <ul style="list-style-type: none"> temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: <ul style="list-style-type: none"> nasal congestion sneezing (Nighttime only) runny nose (Nighttime only) itchy, watery eyes (Nighttime only) itching of the nose or throat (Nighttime only) temporarily relieves these symptoms due to the common cold: <ul style="list-style-type: none"> nasal congestion sneezing (Nighttime only) runny nose (Nighttime only) temporarily relieves sinus congestion and pressure 	<p>Drug Facts (continued)</p> <p>Warnings</p> <p>Do not use:</p> <ul style="list-style-type: none"> 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. with any other product containing diphenhydramine, even one used on skin (Nighttime only) <p>Ask a doctor before use if you have:</p> <ul style="list-style-type: none"> heart disease diabetes thyroid disease high blood pressure difficulty in urination due to enlargement of the prostate gland a breathing problem such as emphysema or chronic bronchitis (Nighttime only) glaucoma (Nighttime only) <p>Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers. (Nighttime only)</p>
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FBI PEEL HERE FOR MORE DRUG FACTS

SINUS PRESSURE & CONGESTION RELIEF PE

PHENYLEPHRINE HCl DIPHENHYDRAMINE HCl

NDC 11822-9485-9 Compare to the active ingredients in Sudafed PE® Sinus Congestion Day + Night®

SINUS PRESSURE & CONGESTION RELIEF PE

<p>DAYTIME</p> <p>PHENYLEPHRINE HCl NASAL DECONGESTANT</p> <p>NON-DROWSY</p> <p>Relieves nasal congestion</p> <p>12 DAYTIME TABLETS</p>	<p>NIGHTTIME</p> <p>DIPHENHYDRAMINE HCl PHENYLEPHRINE HCl ANTIHISTAMINE, NASAL DECONGESTANT</p> <p>Relieves nasal congestion, runny nose</p> <p>8 NIGHTTIME TABLETS</p>
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SINUS PRESSURE & CONGESTION RELIEF PE

PHENYLEPHRINE HCl PHENYLEPHRINE HCl

SATISFACTION GUARANTEE

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

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

*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark SUDAFED PE® SINUS CONGESTION DAY + NIGHT.

50844 REV0820045348509

Do Not Take Daytime and Nighttime Products at the Same Time.

Rite Aid 44-453485

SINUS PRESSURE AND CONGESTION RELIEF

diphenhydramine hcl, phenylephrine hcl kit

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-9485
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-9485-9	1 in 1 CARTON; Type 0: Not a Combination Product	07/02/2019	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BLISTER PACK	12
Part 2	1 BLISTER PACK	8

Part 1 of 2

DAYTIME SINUS PRESSURE AND CONGESTION RELIEF

phenylephrine hcl tablet, film coated

Product Information

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		12 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	07/02/2019	

Part 2 of 2

NIGHTTIME SINUS PRESSURE AND CONGESTION RELIEF

diphenhydramine hcl, phenylephrine hcl tablet, film coated

Product Information

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII: 8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
DIBASIC CALCIUM PHOSPHATE DIHYDRATE (UNII: O7TSZ97GEP)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	blue	Score	no score
Shape	OVAL	Size	11mm
Flavor		Imprint Code	44;485
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		8 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		07/02/2019	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012		07/02/2019	

Labeler - Rite Aid Corporation (014578892)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(11822-9485) , pack(11822-9485)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(11822-9485)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(11822-9485)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(11822-9485)

Revised: 7/2023

Rite Aid Corporation