

**SUPHEDRINE PE SINUS CONGESTION NON-DROWSY- phenylephrine hcl tablet,
film coated
ARMY AND AIR FORCE EXCHANGE SERVICE**

Exchange Select 44-453-Sinus

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

exchange✓ **select**™

Compare To The Active Ingredient of SUDAFED PE® SINUS CONGESTION*

**SUPHEDRINE PE
SINUS CONGESTION**

Phenylephrine HCl 10 mg, Nasal Decongestant

- Maximum Strength
- Sinus Pressure
- Congestion

NON-DROWSY

Actual Size

72 Tablets

quality value

**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 REV0820L45323

"SATISFACTION GUARANTEED OR YOUR MONEY BACK"

Manufactured For Your Military Exchanges

Distributed by: LNK International, Inc.

Hauppauge, NY 11788

1-800-426-9391

SUPHEDRINE PE

SINUS CONGESTION

exchange **select**

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No Print Area
Lot & Exp Date

SINUS CONGESTION
SUPHEDRINE PE

B-0066F-453-23-R
REV0820L45323



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SINUS CONGESTION, 50844 REV0820L45223
"SATISFACTION GUARANTEED OR YOUR MONEY BACK"
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Drug Facts	Other Information	Questions or comments? 1-800-426-9391
<p>Drug Facts (continued)</p> <p>Active ingredient (in each tablet) Phenylephrine HCl 10 mg Nasal decongestant</p> <p>Uses temporarily relieves sinus congestion and pressure temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies</p> <p>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.</p> <p>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</p> <p>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</p>	<p>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</p> <p>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</p> <p>Inactive ingredients croscarmellose sodium, dextrose monohydrate, dibasic calcium phosphate dihydrate, F&C red #40, lecithin, magnesium stearate, maldextrin, microcrystalline cellulose, silicon dioxide, sodium carboxymethylcellulose, sodium citrate dihydrate, titanium dioxide</p>	<p>Questions or comments? 1-800-426-9391</p>

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

44-453

SUPHEDRINE PE SINUS CONGESTION NON-DROWSY

phenylephrine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55301-753
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
CROSCARMELLOSE 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: 70097M6130)	
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:55301-753-23	3 in 1 CARTON	01/15/2021	
1		24 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:55301-753-44	1 in 1 CARTON	01/15/2021	
2		18 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
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Category	Citation	Date	Date
OTC Monograph Drug	M012	01/15/2021	

Labeler - ARMY AND AIR FORCE EXCHANGE SERVICE (001695568)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(55301-753) , pack(55301-753)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(55301-753)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(55301-753)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(55301-753)

Revised: 1/2024

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