

SINUS CONGESTION- phenylephrine hcl tablet, film coated
L.N.K. International, Inc.

Quality Plus 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

**QUALITY
+PLUS**

NDC 50844-493-08

*Compare to active ingredient in Sudafed PE® Sinus Congestion

**MAXIMUM STRENGTH
SINUS CONGESTION**

Phenylephrine HCl 10 mg • NASAL DECONGESTANT

- SINUS PRESSURE
- SINUS CONGESTION

24 Tablets

NON-DROWSY

ACTUAL SIZE

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 ORG082045308

Distributed by

LNK INTERNATIONAL, INC.

60 Arkay Drive, Hauppauge, NY 11788
USA

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.

Directions

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Other information

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Questions or comments? 1-800-426-9391

MAXIMUM STRENGTH

SINUS CONGESTION

3-1603-453-08-R
ORG082045308

NDC 50844-493-08

*Compare to active ingredient in Sudafed PE® Sinus Congestion

MAXIMUM STRENGTH

SINUS CONGESTION

Phenylephrine HCl 10 mg • NASAL DECONGESTANT

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SINUS CONGESTION • SINUS PRESSURE • SINUS CONGESTION • SINUS

24 Tablets • SINUS NON-DROWSY PRESSURE • ACTUAL SIZE

QUALITY  PLUS
SINUS CONGESTION
MAXIMUM STRENGTH

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

No Print / No Varnish Lot no. & Exp. date

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60 Arkay Drive, Hauppauge, NY 11788
USA



Quality Plus 44-453

SINUS CONGESTION

phenylephrine hcl tablet, film coated

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:50844-493
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: 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:50844-493-08	1 in 1 CARTON	07/14/2021	
1		24 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:50844-493-07	2 in 1 CARTON	05/17/2023	
2		18 in 1 BLISTER PACK; Type 0: Not a Combination Product		
Marketing Information				
Marketing		Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M012	07/14/2021	

Labeler - L.N.K. International, Inc. (038154464)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(50844-493) , pack(50844-493)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(50844-493)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(50844-493)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(50844-493)