

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

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**Sound Body 44-453-Sinus Congestion**

***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***

SOUNDBODY™

\*Compare to the active ingredient  
in SUDAFED PE® SINUS CONGESTION

NDC 50844-443-07

MAXIMUM STRENGTH

***Sinus Congestion PE***

**Phenylephrine HCl 10 mg**, Nasal Decongestant

Relieves Sinus Pressure and Congestion

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 Johnson &

Johnson Corporation, owner of the registered trademark SUDAFED PE®  
SINUS CONGESTION. 50844 ORG082045307

Manufactured for Big Lots Stores, Inc.

by **LNK INTERNATIONAL, INC.**

60 Arkay Drive, Hauppauge, NY 11788 USA

V#733000 ITEM#022745307BLBX



MAXIMUM STRENGTH

# Sinus Congestion PE

Phenylephrine HCl 10 mg, Nasal Decongestant



MAXIMUM STRENGTH  
**Sinus Congestion PE**  
Phenylephrine HCl 10 mg, Nasal Decongestant



\*Compare to the active ingredient  
in SUDAFED PE® SINUS CONGESTION

NDC 50844-443-07

MAXIMUM STRENGTH

# Sinus Congestion PE

Phenylephrine HCl 10 mg, Nasal Decongestant  
Relieves Sinus Pressure and Congestion

Non Drowsy

**36** TABLETS

Actual Size



No Print/No varnish Area  
Lot and Exp Date

B-0227-453-07-RR  
OR6082045307

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SINUS CONGESTION. 50844 OR6082045307  
Manufactured for Big Lots Stores, Inc.  
by LNK INTERNATIONAL, INC.  
60 Arkay Drive, Haverhill, MA 01830 USA  
V#733000 ITEM#022745307BLB

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION	
<b>Drug Facts</b>	<b>Active ingredient (in each tablet)</b> Phenylephrine HCl 10 mg ..... Nasal decongestant
<b>Uses</b>	■ temporarily relieves nasal congestion due to the common cold, ■ temporarily relieves sinus congestion and pressure
<b>Warnings</b>	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
<b>When using this product do not exceed recommended dosage.</b>	■ nervousness, dizziness, or sleeplessness occur ■ symptoms do not improve within 7 days or occur with fever
<b>Stop use and ask a doctor if</b>	
<b>Other information</b>	■ see end flap for expiration date and lot number ■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F) ■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN ■ 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
<b>Directions</b>	Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.
<b>Drug Facts (continued)</b>	<b>Ingredients</b> Inactive ingredients: croscarmellose sodium, dextrose monohydrate, dibasic calcium phosphate dihydrate, FD&C red #40, lecithin, magnesium stearate, maito dextrin, microcrystalline cellulose, silicon dioxide, sodium carboxymethylcellulose, sodium citrate dihydrate, titanium dioxide
<b>Questions or comments?</b>	1-800-426-9391

Sound Body 44-453

## SINUS CONGESTION PE

phenylephrine hcl tablet, film coated

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50844-443	
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:50844-443-07	2 in 1 CARTON	12/01/2020	
1		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	12/01/2020	

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

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(50844-443) , pack(50844-443)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(50844-443)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(50844-443)

## Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(50844-443)

Revised: 12/2023

L.N.K. International, Inc.