# NASAL DECONGESTANT- phenylephrine hcl tablet, film coated L.N.K. International, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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#### Convenience Valet 44-453-Delisted

#### 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 (1-800-222-1222) 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?

1-844-428-2538

#### Principal display panel

24/7 BY **7-Eleven**™

Non-Drowsy Maximum Strength Nasal Decongestant Phenylephrine HCl/Nasal Decongestant

**Relieves:** Sinus Pressure, Sinus & Nasal Congestion

compare to Sudafed PE® Congestion active ingredient\* 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® Congestion. 50844 ORG011845344

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#### Satisfaction Guaranteed 1-800-255-0711



# NASAL DECONGESTANT

phenylephrine hcl tablet, film coated

Product Type Route of Administ Active Ingredier PHENYLEPHRINE HYD UNII: 1WS 297W6MV) Inactive Ingredi CROSCARMELLOSE S DEXTROSE MONOHY DIBASIC CALCIUM PH FD&C RED NO. 40 (U LECITHIN, SOYBEAN MAGNESIUM STEARA MALTODEXTRIN (UNII MICROCRYSTALLINE SILICON DIOXIDE (UN CARBOXYMETHYLCE TRISODIUM CITRATE TITANIUM DIOXIDE (UN	nt/Active Ingre DROCHLORI ients ients SODIUM (UN 'DRATE (UNI 'DRATE (UNI UNII: 1DI56 ATE (UNII: 1DI56 ATE (UNII: 70 I: 7CVR7L4A2 CELLULOSI WII: ETJ7Z 6XE ELLULOSE SE DIHYDRATI	Ingredie           IDE (UNII: 04JA59           IDIHYDRATE (UN           7XOA)           QDM62)           097M6I30)           2D)           E (UNII: 041R32E           304)           ODIUM, UNSPE           E (UNII: 8225478	TNSJ) (PHENYLEI nt Name II: 07TSZ97GEP 061U) CIFIED (UNII: K6	)	Basis of St PHENYLEPHRINE HYDROCHLORIDE	=	44-263 Strengtl 10 mg
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TRISODIUM CITRATE	DIHYDRAT	<b>E</b> (UNII: B22547B		790BS311	)		
			95K)				
TITANIUM DIOXIDE (I	UNII: 15FIX9V	/2IP)					
Product Charact	teristics						
Color	red		Score		1	no score	
Shape		JND	Size			7mm	
Flavor			Imprint Code			44;453	
Contains						11,155	
contains							
Packaging							
# Item Code	Ра	ickage Descr	iption	Mar	keting Start Date		ting End ate
<b>1</b> NDC:50844- 263-44 1 i	in 1 CARTON			08/05/	2019	06/16/202	4
	3 in 1 BLISTE oduct	R PACK; Type 0:	Not a Combinati	on			
Marketing In	format	ion					
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OTC monograph final	part341			08/05/	/2019	06/16/202	24

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

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		038154464	pack(50844-263)
Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		832867837	pack(50844-263)
Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		832867894	manufacture(50844-263)
Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		868734088	pack(50844-263)
Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		967626305	pack(50844-263)

Revised: 6/2022

L.N.K. International, Inc.