

SINUS CONGESTION- phenylephrine hcl tablet, film coated
Kroger Company

Kroger 44-453-Sinus Congestion

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?

1-800-632-6900

Principal display panel

Kroger®

COMPARE TO the active ingredient of SUDAFED PE® CONGESTION *See bottom panel

NDC 30142-853-07

Maximum Strength

Sinus Congestion

Phenylephrine HCl 10 mg

Nasal Decongestant

NON-DROWSY

RELIEVES:

- Sinus Pressure
- Nasal & Sinus Congestion

actual size

36 TABLETS

Our Pharmacists Recommended

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

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www.kroger.com**

**DISTRIBUTED BY THE KROGER CO.
CINCINNATI, OHIO 45202**

**FOR MORE PRODUCT INFORMATION, SCAN UPC
USING YOUR **KROGER APP** OR CALL 800-632-6900**

*Sudafed PE® Sinus Congestion is a registered trademark of Johnson & Johnson Corporation, New Brunswick, NJ 08933. Johnson & Johnson Corporation is not affiliated with the Kroger Co. or this product.

50844 ORG082045307



Maximum Strength

Sinus Congestion

COMPARE TO the active ingredient of SUDAFED PE® SINUS CONGESTION *See bottom panel

NDC 30142-853-07



Maximum Strength

Sinus Congestion

Phenylephrine HCl 10 mg
Nasal Decongestant

NON-DROWSY

RELIEVES:

Sinus Pressure
Nasal & Sinus Congestion

actual
size



36 TABLETS



85% SIZE
12-digit UPC-A
(non-suppressed)
For Position Only

0 41260 01216 7

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



PLEASE
RECYCLE

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50844 ORG082045307

B-1050-453-07-RR
ORG082045307

Drug Facts Drug Facts (continued)	Drug Facts Phenylephrine HCl 10 mg Nasal decongestant
Purpose 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.	Uses ■ temporarily relieves sinus congestion and pressure ■ temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies
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	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 ■ symptoms do not improve within 7 days or occur with fever ■ nervousness, dizziness, or sleeplessness occur
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	Active ingredient (in each tablet) ■ symptoms do not improve within 7 days or occur with fever ■ nervousness, dizziness, or sleeplessness occur
Inactive ingredients croscarmellose sodium, dextrose monohydrate, dibasic calcium phosphate dihydrate, FD&C red #40, lecithin, magnesium stearate, maltodextrin, microcrystalline cellulose, silicon dioxide, sodium dodecyl sulfate, titanium dioxide	Questions or comments? 1-800-632-6900

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

Kroger 44-453

SINUS CONGESTION
phenylephrine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30142-853
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	NDC:30142-853-07	2 in 1 CARTON	03/19/2021	
1		18 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:30142-853-23	3 in 1 CARTON	03/19/2021	
2		24 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	03/19/2021	

Labeler - Kroger Company (006999528)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(30142-853) , pack(30142-853)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(30142-853)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(30142-853)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(30142-853)

Revised: 3/2025

Kroger Company