

**SINUS CONGESTION PE- phenylephrine hcl tablet, film coated
WALMART INC.**

Equate 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?

1-888-287-1915

Principal display panel

equate[™]

NDC 79903-076-23

**Compare
to SUDAFED
PE® SINUS
CONGESTION
active
ingredient***

NON-DROWSY

Sinus Congestion PE

Phenylephrine HCl 10 mg

Nasal Decongestant

MAXIMUM STRENGTH

Relieves:

- Sinus Pressure
- Sinus & Nasal Congestion

Actual Size

10

mg
EACH

24
TABLETS

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

Satisfaction guaranteed - Or we'll replace it or give
you your money back. For questions or comments
please call **1-888-287-1915**.

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716

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owner of the registered trademark SUDAFED PE® SINUS CONGESTION.

50844 ORG082045308



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or comments or to report an
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No Print/ No Varnish
Lot #/ Exp. Date

Drug Facts
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

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

When using this product do not exceed recommended dosage.
difficulty in urination due to enlargement of the prostate gland

Stop use and ask a doctor if
■ nervousness, dizziness, or sleepiness occur
■ symptoms do not improve within 7 days or occur with fever

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
■ 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? 1-888-287-1915

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION



equate™

NON-DROWSY

Sinus Congestion PE
Phenylephrine HCl 10 mg

B-2203-453-08
ORG082045308

equate™

NON-DROWSY

Sinus Congestion PE

Phenylephrine HCl 10 mg
Nasal Decongestant

MAXIMUM STRENGTH

Relieves:
• Sinus Pressure
• Sinus & Nasal Congestion



10 mg EACH

24 TABLETS

Actual Size

Equate 44-453

SINUS CONGESTION PE

phenylephrine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-076
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:79903-076-24	1 in 1 CARTON	08/31/2021	03/31/2026
1	NDC:79903-076-43	24 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:79903-076-23	3 in 1 CARTON	08/31/2021	03/31/2026
2	NDC:79903-076-43	24 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	08/31/2021	03/31/2026

Labeler - WALMART INC. (051957769)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(79903-076) , pack(79903-076)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(79903-076)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(79903-076)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(79903-076)

Revised: 7/2025

WALMART INC.