

SINUS AND ALLERGY RELIEF PE- chlorpheniramine maleate, phenylephrine hcl tablet

Topco Associates, LLC

TopCare 44-462

Active ingredients (in each tablet)

Chlorpheniramine maleate 4 mg

Phenylephrine HCl 10 mg

Purpose

Antihistamine

Nasal decongestant

Uses

- temporarily relieves these symptoms due to hay fever (allergic rhinitis) or other upper respiratory allergies:
 - runny nose
 - sneezing
 - itching of the nose or throat
 - itchy, watery eyes
 - sinus congestion and pressure
 - nasal congestion

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

- high blood pressure
- heart disease
- thyroid disease
- diabetes
- glaucoma
- difficulty in urination due to enlargement of the prostate gland
- a breathing problem such as emphysema or chronic bronchitis

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- **do not exceed recommended dosage**
- excitability may occur, especially in children
- drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives and tranquilizers may increase drowsiness
- use caution when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- symptoms do not improve within 7 days or occur with a 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: do not use

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, lactose anhydrous, magnesium stearate, microcrystalline cellulose, silicon dioxide, stearic acid

Questions or comments?

1-888-423-0139

Principal display panel

+TopCare®

health

NDC 36800-642-08

MAXIMUM STRENGTH

Sinus & Allergy Relief PE

CHLORPHENIRAMINE MALEATE 4 mg - ANTIHISTAMINE
PHENYLEPHRINE HCl 10 mg - NASAL DECONGESTANT

RELIEVES:

- Sneezing
- Runny Nose
- Nasal Congestion
- Sinus Congestion & Pressure
- Itching of the Nose or Throat
- Itchy, Watery Eyes

24 TABLETS

actual size

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

DISTRIBUTED BY TOPCO ASSOCIATES LLC

ELK GROVE VILLAGE, IL 60007

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QUESTIONS? 1-888-423-0139

topcare@topco.com

www.topcarebrand.com

50844 ORG031946208

QUALITY GUARANTEED

This TopCare® product is laboratory tested to guarantee its highest quality. Your total satisfaction is guaranteed.



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No print/No varnish
Lot & Exp date

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Sinus & Allergy Relief PE

CHLORPHENIRAMINE MALEATE 4 mg - ANTIHISTAMINE
PHENYLEPHRINE HCl 10 mg - NASAL DECONGESTANT

B-1910-462-08-R
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Scan here for more information or call 1-888-423-0139



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Drug Facts (continued)	Drug Facts
Purpose Chlorpheniramine maleate 4 mg, Antihistamine Phenylephrine HCl 10 mg, Nasal decongestant	Uses temporarily relieves these symptoms due to hay fever (allergic rhinitis) or other upper respiratory allergies: runny nose, sneezing, itching of the nose or throat, itchy, watery eyes sinus congestion and pressure, nasal congestion
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 high blood pressure, heart disease, thyroid disease, diabetes, glaucoma, difficulty in urination due to enlargement of the prostate gland, or a breathing problem such as emphysema or chronic bronchitis. Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers.	When using this product do not exceed recommended dosage excitability may occur, especially in children
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: do not use	Other information TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN see end flap for expiration date and lot number store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
Inactive ingredients croscarmellose sodium, magnesium stearate, croscarmellose sodium, microcrystalline cellulose, silicon dioxide, stearic acid, lactose anhydrous, magnesium stearate, croscarmellose sodium, lactose anhydrous, croscarmellose sodium, microcrystalline cellulose, silicon dioxide, stearic acid	Questions or comments? 1-888-423-0139

TopCare 44-462

SINUS AND ALLERGY RELIEF PE

chlorpheniramine maleate, phenylephrine hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:36800-642
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	4 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	7mm
Flavor		Imprint Code	44;462
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-642-08	1 in 1 CARTON	07/02/2021	
1		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	07/02/2021	

Labeler - Topco Associates, LLC (006935977)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(36800-642) , pack(36800-642)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(36800-642)

Revised: 7/2025

Topco Associates, LLC