SINUS AND ALLERGY MAXIMUM STRENGTH- chlorpheniramine maleate, phenylephrine hcl tablet
Chain Drug Consortium

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.

Premier Value 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
- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers.

When using this product

- **do not exceed recommended dose**
- excitability may occur, especially in children
- drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives and tranquilizers may increase drowsiness
- be careful 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 this product in children under 12 years of age

Other information
- 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, magnesium stearate, microcrystalline cellulose, silica gel, stearic acid

Questions or comments?

1-800-426-9391

Principal Display Panel

Premier
***Value*®**

NDC 68016-687-24

***COMPARE TO THE ACITVE INGREDIENTS IN**
SUDAFED PE® SINUS + ALLERGY

Maximum Strength
Sinus & Allergy

Chlorpheniramine maleate 4 mg - **ANTIHISTAMINE**
Phenylephrine HCl 10 mg - **NASAL DECONGESTANT**

• Sneezing • Itchy eyes • Runny nose
• Sinus pressure + congestion

24 Tablets

INDEPENDENTLY TESTED
PV
SATISFACTION GUARANTEED

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

*This product is not manufactured or distributed by McNeil Consumer Healthcare, owner of the registered trademark Sudafed PE® Sinus + Allergy.
50844 ORG061246208

DISTRIBUTED BY: CHAIN CONSORTIUM, LLC
UPARC, BLDG. A3, SUITE 338
1020 WILLIAM PITT WAY
PITTSBURGH, PA 15238
www.chaindrugconsortium.com

If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.

Premier Value 44-462

SINUS AND ALLERGY MAXIMUM STRENGTH

chlorpheniramine maleate, phenylephrine hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-687
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
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

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:68016-687-24	1 in 1 CARTON	06/09/2005	
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 FINAL	part341	06/09/2005	

Labeler - Chain Drug Consortium (101668460)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	MANUFACTURE(68016-687)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	PACK(68016-687)

Revised: 6/2019

Chain Drug Consortium