

**SINUS PLUS ALLERGY PE MAXIMUM STRENGTH- chlorpheniramine maleate,
phenylephrine hcl tablet
United Natural Foods, Inc. dba UNFI**

Equaline 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
- difficulty in urination due to enlargement of the prostate gland

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°C-30°C (59°F-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?

1-855-423-2630

Principal Display Panel

EQUALINE®

NDC 41163-666-08

maximum strength
sinus + allergy PE

chlorpheniramine maleate 4 mg
(antihistamine)
phenylephrine HCl 10 mg
(nasal decongestant)

relieves:

- *sneezing, itchy eyes, runny nose*
- *sinus pressure & congestion*

24 tablets

PSEUDOEPHEDRINE FREE

actual size

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

50844

REV0319B46208

**DISTRIBUTED BY UNFI
PROVIDENCE, RI 02908 USA
855-423-2630**

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maximum strength
sinus + allergy PE

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TAMPER EVIDENT: DO NOT USE IF
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No print area
Lot no/Exp date

maximum strength
sinus + allergy PE

EQUALINE®

DISTRIBUTED BY UNFI
PROVIDENCE, RI 02908 USA
855-423-2630

**100% Quality
GUARANTEED**

50844 REV0319B46208

B-0019E-462-08-R
REV0319B46208

Drug Facts	
KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION	
Active ingredients (in each tablet) 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: ■ itchy, watery eyes ■ runny nose ■ itching of the nose or throat ■ nasal congestion ■ sneezing ■ 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 ■ 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. Do not exceed recommended dosage ■ do not exceed recommended dosage ■ excitability may occur, especially in children ■ drowsiness may occur	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	Drug Facts (continued) 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 ■ symptoms do not improve within 7 days or occur with a fever ■ nervousness, dizziness, or sleepiness occur 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.
Inactive ingredients croscarmellose sodium, lactose anhydrous, magnesium stearate, microcrystalline cellulose, silicon dioxide, stearic acid	Questions or comments? 1-855-423-2630

Equaline 44-462

SINUS PLUS ALLERGY PE MAXIMUM STRENGTH

chlorpheniramine maleate, phenylephrine hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41163-666
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:41163-666-08	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 Drug	M012	06/09/2005	

Labeler - United Natural Foods, Inc. dba UNFI (943556183)

Establishment

Name	Address	ID/FEI	Business Operations
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LNK International, Inc.		832867837	manufacture(41163-666) , pack(41163-666)
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Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(41163-666)

Revised: 1/2024

United Natural Foods, Inc. dba UNFI