

NASAL DECONGESTANT PE- phenylephrine hcl tablet, film coated
United Natural Foods, Inc. dba UNFI

Equaline 44-453-Delisted

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?

1-855-423-2630

Principal Display Panel

EQUALINE®

NDC 41163-453-07

**compare to
Sudafed PE® Sinus
Congestion**

active ingredient*

maximum strength

nasal

decongestant PE

phenylephrine HCl 10 mg (nasal decongestant)

non-drowsy

relieves:

- *sinus pressure*
- *congestion*

36 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**

**100% *Quality*
GUARANTEED**

Like it or let us
make it right.

**That's our
quality promise.**

855-423-2630

**DISTRIBUTED BY UNFI
PROVIDENCE, RI 02908 USA**

*This product is not manufactured or
distributed by Johnson & Johnson
Corporation, owner of the registered
trademark Sudafed PE® Sinus
Congestion.

50844 REV0820E45307

EQUALINE®

maximum strength
nasal decongestant PE

NDC 41163-453-07

compare to
**Sudafed PE® Sinus
Congestion**
active ingredient*

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

maximum strength
nasal decongestant PE

EQUALINE®

maximum strength
**nasal
decongestant PE**

phenylephrine HCl 10 mg (nasal decongestant)

non - drowsy

relieves:
• sinus pressure
• congestion

36 tablets

actual size

No Print Area
Lot & Exp

B-0019E-453-07-R
REV0820E45307



*This product is not manufactured or
distributed by Johnson & Johnson
Corporation, owner of the registered
trademark Sudafed PE® Sinus
Congestion.
50844 REV0820E45307

Like it or let us
make it right.
That's our
quality promise.
855-425-2630



DISTRIBUTED BY UNFI
PROVIDENCE, RI 02908 USA

Drug Facts	Active ingredient (in each tablet) Phenylephrine HCl 10 mg Nasal decongestant
Purpose	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.
Uses	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
Directions	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
Other information	■ see end flap for expiration date and lot number ■ store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F) ■ TAMPER EVIDENT: DO NOT USE IF OUTER PACKAGE IS OPENED OR BLISTER IS TORN OR BROKEN
Drug Facts (continued)	Drug Facts 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.
Questions or comments?	1-855-423-2630

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

Equaline 44-453

NASAL DECONGESTANT PE
phenylephrine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41163-453
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:41163-453-07	2 in 1 CARTON	01/14/2005	03/31/2026
1		18 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	01/14/2005	03/31/2026

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

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(41163-453) , pack(41163-453)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(41163-453)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		868734088	manufacture(41163-453)

Establishment

Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(41163-453)

Revised: 8/2025

United Natural Foods, Inc. dba UNFI