

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

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**Equaline 44-453**

***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  
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TAMPERING

maximum strength  
**nasal decongestant PE**

**EQUALINE®**

**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



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Corporation, owner of the registered  
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Congestion.  
50844 REV0820E45307

DISTRIBUTED BY UNFI  
PROVIDENCE, RI 02908 USA

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



<b>Drug Facts</b>	<b>Active ingredient (in each tablet)</b> Phenylephrine HCl 10 mg ..... Nasal decongestant
<b>Purpose</b>	temporarily relieves sinus congestion and pressure due to the common cold, hay fever or other upper respiratory allergies
<b>Warnings</b>	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.
<b>Uses</b>	Ask a doctor before use if you have heart disease ■ diabetes ■ high blood pressure ■ thyroid disease ■ difficulty in urination due to enlargement of the prostate gland
<b>Directions</b>	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
<b>Other information</b>	■ 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
<b>Drug Facts (continued)</b>	<b>Questions or comments?</b> 1-855-423-2630
<b>Drug Facts (continued)</b>	<b>Inactive ingredients</b> croscarmellose sodium, dextrose monohydrate, dibasic calcium phosphate dihydrate, FD&C red #40, lecithin, magnesium stearate, malto-dextrin, microcrystalline cellulose, silicon dioxide, sodium carboxymethylcellulose, sodium citrate dihydrate, titanium dioxide

KEEP OUTER PACKAGE FOR COMPLETE PRODUCT INFORMATION

**Equaline 44-453**

**NASAL DECONGESTANT PE**

phenylephrine hcl tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:41163-453
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>DEXTROSE MONOHYDRATE</b> (UNII: LX22YL083G)	
<b>DIBASIC CALCIUM PHOSPHATE DIHYDRATE</b> (UNII: O7TSZ97GEP)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>LECITHIN, SOYBEAN</b> (UNII: 1DI56QDM62)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED</b> (UNII: K679OBS311)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

**Product Characteristics**

<b>Color</b>	red	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	7mm
<b>Flavor</b>		<b>Imprint Code</b>	44;453
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-453-07	2 in 1 CARTON	01/14/2005	
1		18 in 1 BLISTER PACK; Type 0: Not a Combination Product		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug	M012	01/14/2005	

**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: 12/2023

United Natural Foods, Inc. dba UNFI