

**SINUS CONGESTION AND PAIN DAYTIME- acetaminophen and phenylephrine
hcl tablet, film coated
United Natural Foods, Inc. dba UNFI**

Equaline 44-466C

Active ingredients (in each caplet)

Acetaminophen 325 mg
Phenylephrine HCl 5 mg

Purpose

Pain reliever/fever reducer
Nasal decongestant

Uses

- temporarily relieves these symptoms associated with hay fever or other respiratory allergies, and the common cold:
 - minor aches and pains
 - headache
 - nasal congestion
 - sinus congestion and pressure
- helps decongest sinus openings and passages
- promotes sinus drainage
- helps clear nasal passages
- temporarily reduces fever

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- blisters
- rash
- skin reddening

If a skin reaction occurs, stop use and seek medical help right away.

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If

you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

- 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.
- if you have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if you have

- difficulty in urination due to enlargement of the prostate gland
- liver disease
- heart disease
- diabetes
- thyroid disease
- high blood pressure

Ask a doctor or pharmacist before use if you are

taking the blood thinning drug warfarin.

When using this product

do not exceed recommended dosage.

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- pain or nasal congestion gets worse or lasts more than 7 days
- new symptoms occur
- fever gets worse or lasts more than 3 days
- redness or swelling is present

These could be signs of a serious condition.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of accidental overdose, get medical help or contact a Poison Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- **do not take more than directed**
- adults and children 12 years and over
 - take 2 caplets every 4 hours
 - swallow whole - do not crush, chew, or dissolve
 - do not take more than 10 caplets 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

corn starch, crospovidone, D&C yellow #10 aluminum lake, FD&C blue #1 aluminum lake, FD&C red #40 aluminum lake, flavor, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, sucralose, talc, titanium dioxide

Questions or comments?

1-855-423-2630

Principal Display Panel

NDC 41163-664-08

EQUALINE®

compare to

Tylenol®

SINUS + HEADACHE

active ingredients*

**daytime sinus
congestion & pain
acetaminophen**

(pain reliever/fever reducer)

phenylephrine HCl

(nasal decongestant)

non-drowsy

relieves:

- *sinus headache & pressure*
- *nasal congestion*

PSEUDOEPHEDRINE FREE

24 caplets

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**

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

*This product is not manufactured or distributed by
Johnson & Johnson Corporation, owner of the
registered trademark Tylenol® SINUS + HEADACHE.

50844 REV0721H46608

Drug Facts
 COMPLETE PRODUCT INFORMATION
 KEEP OUTER PACKAGE FOR
Active ingredients
 (in each caplet)
 Acetaminophen 325 mg Pain reliever/fever reducer
 Phenylephrine HCl 5 mg Nasal decongestant

Drug Facts (continued)
Uses
 ■ temporarily relieves these symptoms associated with hay fever or other respiratory allergies, and the common cold: ■ minor aches and pains ■ headache ■ nasal congestion ■ sinus congestion and pressure

compare to
 Tylenol®
 SINUS + HEADACHE
 active ingredients*

NDC 41163-664-08

EQUALINE®

daytime sinus congestion & pain

acetaminophen
 (pain reliever/fever reducer)
 phenylephrine HCl
 (nasal decongestant)

non-drowsy

- relieves:
- sinus headache & pressure
 - nasal congestion

PSEUDOEPHEDRINE FREE



actual size

24 caplets

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No print area
 Lot no. & Exp. Date

Drug Facts (continued)
Inactive ingredients
 com starch, crospovidone, D & C yellow #10 aluminum lake, FD & C blue #1 aluminum lake, FD & C red #40 aluminum lake, flavor, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, povidone,

Drug Facts (continued)
 silicon dioxide, sodium starch glycolate, stearic acid, sucralose, talc, titanium dioxide

Questions or comments? 1-855-423-2630

Do not use
 ■ with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
 ■ 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.
 ■ if you have ever had an allergic reaction to this product or any of its ingredients
 Ask a doctor before use if you have
 ■ difficulty in urination due to enlargement of the prostate gland ■ liver disease ■ heart disease ■ diabetes

Control Center right away. Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions
 ■ do not take more than directed ■ adults and children 12 years and over ■ take 2 caplets every 4 hours ■ swallow whole – do not crush, chew, or dissolve ■ do not take more than 10 caplets in 24 hours ■ children under 12 years: ask a doctor

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100% Quality GUARANTEED
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 855-423-2630

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



No F
 Glue

B-0019E-466C-08SCR
 REV0721H46608

Print Area

Drug Facts (continued)

■ thyroid disease ■ high blood pressure

Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin.

When using this product do not exceed recommended dosage.

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- pain or nasal congestion gets worse or lasts more than 7 days ■ new symptoms occur
- fever gets worse or lasts more than 3 days
- redness or swelling is present

These could be signs of a serious condition.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison

No F Glue

Drug Facts (continued)

- helps decongest sinus openings and passages
- promotes sinus drainage
- helps clear nasal passages
- temporarily reduces fever

Warnings

Liver warning: This product contains acetaminophen. Severe liver damage may occur if you take more than 4,000 mg of acetaminophen in 24 hours

- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

Allergy alert: Acetaminophen may cause severe skin reactions. Symptoms may include:

- hives ■ rash ■ skin reddening

If a skin reaction occurs, stop use and seek medical help right away.

Equaline 44-466C

SINUS CONGESTION AND PAIN DAYTIME

acetaminophen and phenylephrine hcl tablet, film coated

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:41163-664 |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|-----------------------------------------------------------------------------------------|-----------------------------|----------|
| ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) | ACETAMINOPHEN | 325 mg |
| PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 5 mg |

Inactive Ingredients

| Ingredient Name | Strength |
|-----------------------------------------------------------------|----------|
| STARCH, CORN (UNII: O8232NY3SJ) | |
| CROSPVIDONE, UNSPECIFIED (UNII: 2S7830E561) | |
| D&C YELLOW NO. 10 ALUMINUM LAKE (UNII: CQ3XH3DET6) | |
| FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM) | |
| FD&C RED NO. 40 ALUMINUM LAKE (UNII: 6T47AS764T) | |
| MAGNESIUM STEARATE (UNII: 70097M6130) | |
| MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) | |
| POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A) | |
| POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990) | |
| POVIDONE, UNSPECIFIED (UNII: FZ989GH94E) | |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4) | |
| SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2) | |
| STEARIC ACID (UNII: 4ELV7Z65AP) | |
| SUCRALOSE (UNII: 96K6UQ3ZD4) | |
| TALC (UNII: 7SEV7J4R1U) | |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) | |

Product Characteristics

| | | | |
|-----------------|---------|---------------------|----------|
| Color | green | Score | no score |
| Shape | OVAL | Size | 17mm |
| Flavor | MENTHOL | Imprint Code | 44;466 |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---------------------------------------------------------|----------------------|--------------------|
| 1 | NDC:41163-664-08 | 2 in 1 CARTON | 07/26/2005 | |
| 1 | | 12 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 | 07/26/2005 | |

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

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|------------------------------------------|
| LNK International, Inc. | | 832867837 | manufacture(41163-664) , pack(41163-664) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|------------------------|
| LNK International, Inc. | | 832867894 | manufacture(41163-664) |

Establishment

| Name | Address | ID/FEI | Business Operations |
|-------------------------|---------|-----------|------------------------|
| LNK International, Inc. | | 117025878 | manufacture(41163-664) |

Revised: 2/2026

United Natural Foods, Inc. dba UNFI