

COLD AND FLU DAYTIME SEVERE- acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl tablet, film coated
L.N.K. International, Inc.

Convenience Valet 44-640

Active ingredients (in each caplet)

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Guaifenesin 200 mg
Phenylephrine HCl 5 mg

Purpose

Pain reliever/fever reducer
Cough suppressant
Expectorant
Nasal decongestant

Uses

- temporarily relieves common cold and flu symptoms:
 - minor aches and pains
 - sinus congestion and pressure
 - sore throat
 - fever
 - headache
 - nasal congestion
 - cough due to minor throat and bronchial irritation
- reduces swelling of nasal passages
- temporarily restores freer breathing through the nose
- promotes nasal and/or sinus drainage
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

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.

Sore throat warning: If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

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

- liver disease
- thyroid disease
- diabetes
- heart disease
- high blood pressure
- difficulty in urination due to enlargement of the prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema

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, nasal congestion, or cough gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back or occurs with rash or headache that lasts. 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 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**
- do not take more than 8 caplets in 24 hours
- adults and children 12 years and over: take 2 caplets with water every 4 hours
- children under 12 years: ask a doctor

Other information

- **each caplet contains:** sodium 3 mg
- **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, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, polysorbate 80, polyvinyl alcohol, povidone, silicon dioxide, sodium starch glycolate, stearic acid, talc, titanium dioxide

Questions or comments?

1-877-507-6516

Principal display panel

24/7 life

BY **7-ELEVEN™**

Cold & Flu

DayTime Severe

Acetaminophen / Pain Reliever/Fever Reducer

Dextromethorphan HBr / Cough Suppressant

Guaifenesin / Expectorant

Phenylephrine HCl / Nasal Decongestant

Relieves: Headache, Fever, Sore Throat,
Minor Aches & Pains, Chest Congestion,
Thins & Loosens Mucus, Nasal Congestion,
Sinus Pressure, Cough

compare to **Vicks® DayQuil®**

**VapoCOOL™ SEVERE Cold &
Flu + Congestion** active ingredients*

16
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**

PARENTS:
Learn about teen medicine abuse
www.StopMedicineAbuse.org

Distributed by: 7-Eleven, Inc.
Irving, TX 75063 www.7-ELEVEN.com

Satisfaction Guaranteed 1-800-255-0711

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distributed by The Procter & Gamble Company. Vicks®,
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& Gamble Company.

50844

REV0722A64021

65158C-SE-10-22

No Print/No Varnish
Lot # & Exp. Area

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**Cold & Flu
DayTime
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DayQuil®, and VapoCOOL™ are trademarks of The Procter
& Gamble Company.
50844 REV0722A64021 65158C-SE-10-22

Drug Facts (continued)
Inactive ingredients corn starch, croscopolone, FD&C red #40
aluminum lake, FD&C yellow #6 aluminum lake, magnesium stearate,
maltodextrin, microcrystalline cellulose, polyethylene glycol, polyorbate
80, polyvinyl alcohol, povidone, silicon dioxide, sodium starch
glycolate, stearic acid, talc, titanium dioxide
Questions or comments? 1-877-507-6516

65158C-SE-10-22
B-1205-640-21
REV0722A64021

Drug Facts (continued)
persistent or chronic cough such as occurs with smoking, asthma,
chronic bronchitis, or emphysema
**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, nasal congestion, or cough gets worse or lasts more than 7 days
■ fever gets worse or lasts more than 3 days
■ redness or swelling is present ■ new symptoms occur
■ cough comes back or occurs with rash or headache that lasts. These
could be signs of a serious condition.
Keep out of reach of children. In case of 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 with water every
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■ children under 12 years: ask a doctor
Other information ■ each caplet contains: sodium 3 mg
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■ helps loosen phlegm (mucus) and thin bronchial secretions to rid the
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Liver warning: This product contains acetaminophen. Severe liver
damage may occur if you take
■ more than 4,000 mg of acetaminophen in 24 hours
■ 3 or more alcoholic drinks every day while using this product
Allergy alert: Acetaminophen may cause severe skin reactions.
Symptoms may include: ■ rash ■ skin reddening
■ blisters ■ hives ■ severe skin reactions.
Sore throat warning: If sore throat is severe, persists for more than 2
days, is accompanied or followed by fever, headache, rash, nausea, or
vomiting, consult a doctor promptly.
Do not use
■ with any other drug containing acetaminophen (prescription or
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■ 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
■ liver disease ■ thyroid disease ■ diabetes
■ heart disease ■ high blood pressure
■ difficulty in urination due to enlargement of the prostate gland

Convenience Valet 44-640

COLD AND FLU DAYTIME SEVERE

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50844-640
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 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)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
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)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	orange	Score	no score
Shape	OVAL	Size	19mm
Flavor		Imprint Code	44;640
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50844-640-21	2 in 1 CARTON	02/16/2022	
1		8 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	02/16/2022	

Labeler - L.N.K. International, Inc. (038154464)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867837	manufacture(50844-640) , pack(50844-640)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(50844-640)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		117025878	manufacture(50844-640)

Revised: 1/2024 L.N.K. International, Inc.