# COLD AND FLU NIGHTTIME RELIEF- acetaminophen, dextromethorphan hbr, doxylamine succinate solution L.N.K. International, Inc.

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**Convenience Valet 44-014** 

#### Active ingredients (in each 30 mL)

Acetaminophen 650 mg Dextromethorphan HBr 30 mg Doxylamine succinate 12.5 mg

# **Purpose**

Pain reliever/fever reducer Cough suppressant Antihistamine

#### Uses

- temporarily relieves common cold and flu symptoms:
  - runny nose and sneezing
  - sore throat
  - headache
  - minor aches and pains
  - cough due to minor throat and bronchial irritation
- 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:

- skin reddening
- blisters
- rash

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

- difficulty in urination due to enlargement of the prostate gland
- cough that occurs with too much phlegm (mucus)
- liver disease
- glaucoma
- a breathing problem or chronic cough that lasts or 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
- taking sedatives or tranquilizers

# When using this product

- excitability may occur, especially in children
- avoid alcoholic beverages
- marked drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- use caution when driving a motor vehicle or operating machinery

# Stop use and ask a doctor if

- pain 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

- mL = milliliter
- only use the dose cup provided
- do not exceed 4 doses per 24 hours
- adults and children 12 years and over: 30 mL every 6 hours
- children under 12 years: do not use

#### Other information

- each 30 mL contains: sodium 22 mg
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)
- use by expiration date on package

# Inactive ingredients

anhydrous citric acid, D&C yellow #10, FD&C green #3, FD&C yellow #6, flavors, glycerin, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium chloride, sodium citrate dihydrate, sodium saccharin, sucralose

#### Questions or comments?

**1-877-507-6516** (M-F 8AM - 4:30PM CST)

# Principal display panel

**24/7 life** BY **7-ELEVEN**™

Cold & Flu Nighttime Relief

**Acetaminophen** / Pain Reliever/Fever Reducer Dextromethorphan HBr / Cough Suppressant Doxylamine Succinate / Antihistamine

**Relieves:** Headache, Fever, Sore Throat, Minor Aches & Pains, Sneezing, Runny Nose, Cough

Ages 12 Years and Over

compare to VICKS®

NYQUIL® COLD & FLU
active ingredients\*

12 FL OZ (355 mL)

Eucalyptus Mint Flavor

# TAMPER EVIDENT: DO NOT USE IF PRINTED NECK WRAP IS BROKEN OR MISSING

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# Satisfaction Guaranteed 1-800-255-0711

PARENTS:

Learn about teen medicine abuse www.StopMedicineAbuse.org





■ cough comes back or occurs with rash or headache that lasts. These could be signs of a serious condition.

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acetaminophen, dextromethorphan hbr, doxylamine succinate solution

#### **Product Information**

**Product Type** HUMAN OTC DRUG Item Code (Source) NDC:50844-252

Active Ingredient/Active Moiety				
Ingredient Name	<b>Basis of Strength</b>	Strength		
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	650 mg in 30 mL		
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 30 mL		
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 30 mL		

Inactive Ingredients	
Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics				
Color	green	Score		
Shape		Size		
Flavor	MINT (Eucalyptus Mint)	Imprint Code		
Contains				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50844-252- 02	355 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/14/2019	

Marketing Information			
Marketing Category	• • • • • • • • • • • • • • • • • • • •		Marketing End Date
OTC Monograph Drug	M012	07/14/2019	

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

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		967626305	manufacture(50844-252) , pack(50844-252)	

Revised: 7/2023 L.N.K. International, Inc.