# COLD AND FLU DAYTIME RELIEF- acetaminophen, dextromethorphan hbr, phenylephrine hcl solution L.N.K. International, Inc.

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#### **Convenience Valet 44-012**

### Active ingredients (in each 15 mL)

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Phenylephrine HCl 5 mg

# **Purpose**

Pain reliever/fever reducer Cough suppressant Nasal decongestant

#### Uses

- temporarily relieves common cold and flu symptoms:
  - sore throat
  - nasal congestion
  - 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

- adult takes more than 4,000 mg of acetaminophen in 24 hours
- child takes more than 5 doses in 24 hours
- taken with other drugs containing acetaminophen
- adult has 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 the user has

- 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, or emphysema
- heart disease
- diabetes
- thyroid disease
- liver disease
- high blood pressure

# Ask a doctor or pharmacist before use if the user is

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
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- pain, nasal congestion, or cough gets worse or lasts more than 5 days (children) or 7 days (adults)
- 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 exceed 4 doses per 24 hours
- mL = milliliter
- only use the dose cup provided

adults and children 12 years and	30 mL every 4
over	hours
children 6 to under 12 years	15 mL every 4
crilidi en o to under 12 years	hours
children under 6 years	do not use

#### Other information

- each 15 mL contains: sodium 13 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, FD&C yellow #6, flavors, glycerin, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sodium metabisulfite, sodium saccharin, sorbitol, sucralose

#### Questions or comments?

**1-877-507-6516** (8:00AM - 4:30PM CST)

#### Principal display panel

**24/7** *life* by **7-ELEVEN**™

Cold&Flu
DayTime Relief

# For Ages 6 Years & Over

**Acetaminophen** / Pain Reliever/Fever Reducer Dextromethorphan HBr / Cough Suppressant Phenylephrine HCl / Nasal Decongestant

**Relieves:** Minor Aches & Pains, Sore Throat, Fever, Headache, Nasal Congestion, Cough

compare to Vicks®

DayQuil®Cold & Flu
active ingredients\*

Menthol

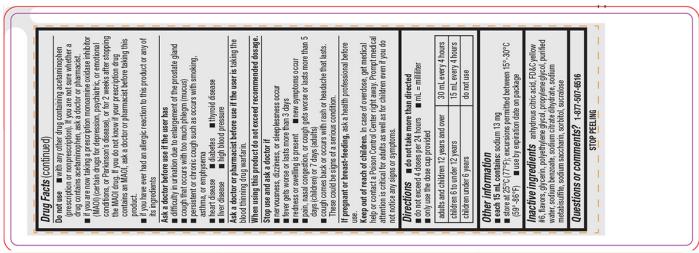
# 6 FL OZ (177 mL)

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

\*This product is not manufactured or distributed by The Procter & Gamble Company, owner of the registered trademark Vicks® DayQuil® Cold & Flu. 50844 ORG032301245

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#### Satisfaction Guaranteed 1-800-255-0711





Convenience valet 44-012

# **COLD AND FLU DAYTIME RELIEF**

acetaminophen, dextromethorphan hbr, phenylephrine hcl solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50844-251
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	325 mg in 15 mL	
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 15 mL	
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 15 mL	

Inactive Ingredients			
Ingredient Name	Strength		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)			
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)			
GLYCERIN (UNII: PDC6A3C0OX)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)			
SODIUM METABISULFITE (UNII: 4VON5FNS3C)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
SORBITOL (UNII: 506T60A25R)			
SUCRALOSE (UNII: 96K6UQ3ZD4)			

Product Characteristics			
Color	orange	Score	
Shape		Size	
Flavor	MENTHOL	Imprint Code	
Contains			

Packaging					
#	Item Code	ode Package Description Marketing Start M Date			
1	NDC:50844- 251-45	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/2019		
2	NDC:50844- 251-02	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/20/2019	04/30/2026	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/20/2019	
OTC Monograph Drug	M012	07/20/2019	

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

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	manufacture(50844-251) , pack(50844-251)

Revised: 8/2025 L.N.K. International, Inc.