# 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

- mL = milliliter
- only use the dose cup provided
- do not exceed 4 doses per 24 hours

adults and children 12 years and	30 mL every 4
over	hours
children 6 to under 12 years	15 mL every 4
children 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 (8:00 AM - 4:30 PM CST)

# Principal display panel

24/7 Life by 7-ELEVEN™

Cold&Flu DayTime Relief

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

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

Ages 6 Years and Over

compare to Vicks® DayQuil® Cold & Flu active ingredients\* Menthol Flavor

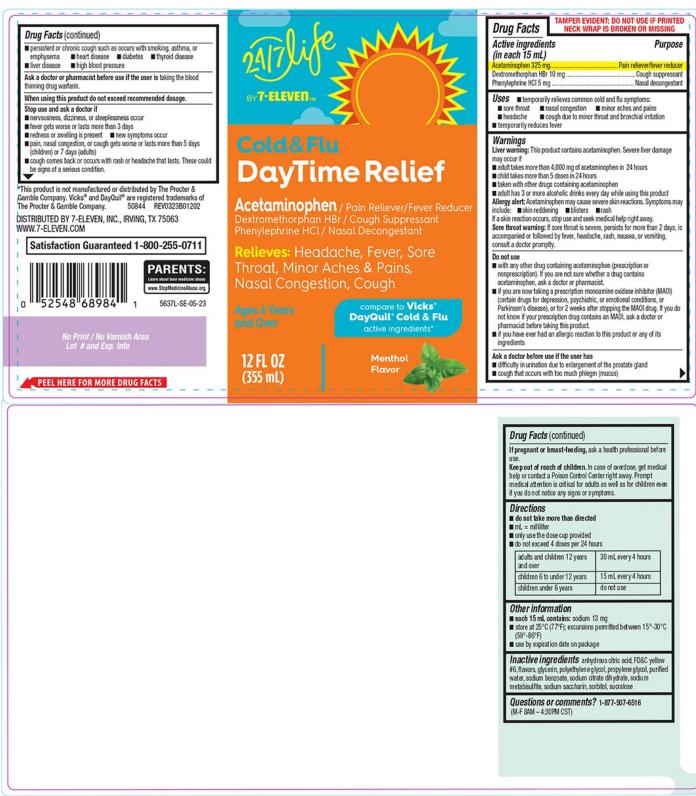
# 12 FL OZ (355 mL)

\*This product is not manufactured or distributed by The Procter & Gamble Company. Vicks ® and DayQuil® are registered trademarks of The Procter & Gamble Company. 50844 REV0323B01202

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# Satisfaction Guaranteed 1-800-255-0711

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



#### **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 Ingred	diant/Activ	e Moietv			
Active ingret				Basis of Stre	ngth Strength
					325 mg
ACETAMINOPHEI	<b>N</b> (UNII: 36209	in 15 mL			
	COMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH)DEXTROMETHORPHANIOMETHORPHAN - UNII:7355X3ROTS)HYDROBROMIDE				
PHENYLEPHRINE UNII:1WS297W6MV	EPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - 297W6MV)PHENYLEPHRINE HYDROCHLORIDE				
Inactive Ingr	edients				
		Ingredient Nam	ne		Strength
ANHYDROUS CIT	RIC ACID (UN	II: XF417D3PSL)			
FD&C YELLOW N	<b>10.6</b> (UNII: H7	7VEI93A8)			
GLYCERIN (UNII: F	PDC6A3C0OX)				
POLYETHYLENE	GLYCOL, UNS	PECIFIED (UNII: 3WQ0SD	WIA)		
PROPYLENE GLY	COL (UNII: 6D	C9Q167V3)			
WATER (UNII: 059	QF0KO0R)				
SODIUM BENZOA	ATE (UNII: OJ24	15FE5EU)			
TRISODIUM CITR	ATE DIHYDRA	<b>TE</b> (UNII: B22547B95K)			
SODIUM METABI	SULFITE (UNI	: 4VON5FNS3C)			
SACCHARIN SOD					
SORBITOL (UNII:					
SUCRALOSE (UNI	•	4)			
		•,			
Product Cha		_			
	racteristic	S			
Color		orange	Score		
Shape		Size			
Flavor		MENTHOL Imprint Co		ode	
Contains					
Packaging					
# Item Code		Package Description		Marketing Start Date	Marketing End Date
<b>1</b> NDC:50844- 251-02	355 mL in 1 E Combination	L BOTTLE, PLASTIC; Type 0: Not a		07/20/2019	
Marketing	Informa	ation			
- Marketing Category		plication Number or Monograph Citation		Marketing Start Date	Marketing End Date
	M012		0.	7/20/2010	

07/20/2019

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

OTC Monograph Drug M012

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

Revised: 7/2023

L.N.K. International, Inc.