#### DAYTIME SEVERE COLD AND FLU- acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl solution Kroger Company

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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# Kroger Co. DayTime Severe Cold & Flu Drug Facts

# Active ingredients (in each 15 mL)

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/flu symptoms:

- sinus congestion and pressure
- nasal congestion
- minor aches and pains
- headache
- sore throat
- fever
- cough due to minor throat and bronchial irritation
- reduces swelling of nasal passages
- promotes nasal and/or sinus drainage
- temporarily restores freer breathing through the nose
- 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

- 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 you have

- liver disease
- heart disease
- thyroid disease
- diabetes
- high blood pressure
- trouble urinating due to an enlarged 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 use more than directed

Stop use and ask a doctor if

- pain, nasal congestion, or cough gets worse or lasts more than 5 days (children) or 7 days (adults)
- you get nervous, dizzy or sleepless
- 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.

**Overdose warning:** In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

# Directions

- take only as directed see Overdose warning
- only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 4 hrs
children 6 to under 12 yrs	15 mL every 4 hrs
children 4 to under 6 yrs	ask a doctor
children under 4 yrs	do not use

# Other information

- each 15 mL contains: sodium 6 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

# Inactive ingredients

butylated hydroxyanisole, edetate disodium, FD&C yellow #6, flavor, glycerin, menthol, monobasic sodium phosphate, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sucrose, xanthan gum

# Questions?

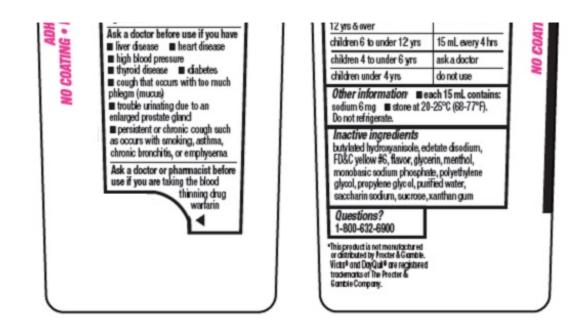
1-800-632-6900

## Package/Label Principal Display Panel

COMPARE TO the active ingredients of VICKS<sup>®</sup> DAYQUIL<sup>®</sup> SEVERE COLD & FLU

See back panel Kroger<sub>®</sub> Maximum Strength DayTime Severe COLD & FLU Acetaminophen Dextromethorphan HBr Phenylephrine HCl Guaifenesin OUR PHARMACIST RECOMENDED Pain Reliever Fever Reducer **Cough Suppressant** Nasal Decongestant Expectorant Non-Drowsy Antihistamine Free Alcohol Free 12 FL OZ (354 mL)





acetaminophen, dextrometho		n, phenyleph	rine hcl solution	1		
Dreduct Information						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:3		NDC:30	80142-603	
Route of Administration	ORAL					
Active Ingredient/Active						
Ingredient Name			<b>Basis of Strength</b>		Strength	
ACETAMINOPHEN (UNII: 36209ITL	9D) (ACETAMINOPHEN - UNII:362O9ITL9D)		ACETAMINOPHEN		325 mg in 15 mL	
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)			DEXTROMETHORPHAN HYDROBROMIDE		10 mg in 15 mL	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)			GUAIFENESIN		200 mg in 15 mL	
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV)			PHENYLEPHRINE HYDROCHLORIDE		5 mg in 15 mL	
Inactive Ingredients						
Ingredient Name					Strength	
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)						
EDETATE DISODIUM (UNII: 7FLD91C86K)						
FD&C YELLOW NO. 6 (UNII: H77V	EI93A8)					
GLYCERIN (UNII: PDC6A3C0OX)						
MENTHOL, UNSPECIFIED FORM						
SODIUM PHOSPHATE, MONOBAS			SW)			
POLYETHYLENE GLYCOL, UNSPE		4)				
PROPYLENE GLYCOL (UNII: 6DC90	Q167V3)					
WATER (UNII: 059QF0KO0R)						
SACCHARIN SODIUM (UNII: SB8Z)	UX40TY)					

	LH8M554)		
XANTHAN GUM (UNII	TIVI2P4NEE)		
Product Charac	teristics		
Color	ORANGE (clear)	Score	
Shape		Size	
Flavor	FRUIT, MENTHOL	Imprint Code	
Contains			
Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
	54 mL in 1 BOTTLE; Type 0: Not a Combination roduct	10/10/2014	
	nformation		
Marketing In			
Marketing Ir Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

# Labeler - Kroger Company (006999528)

Revised: 3/2023

Kroger Company