# COLD AND FLU NIGHTTIME- acetaminophen, dextromethorphan hydrobromide, doxylamine succinate liquid Family Dollar (FAMILY WELLNESS)

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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#### **Drug Facts**

## Active ingredients (in each 15 mL) Acetaminophen 500 mg

Dextromethorphan HBr 15 mg

Doxylamine succinate 6.25 mg

#### **Purposes**

#### Pain reliever/fever reducer

Cough suppressant

**Antihistamine** 

#### Uses

temporarily relieves these common cold/flu symptoms:

- minor aches and pains
- headache
- sore throat
- fever
- runny nose and sneezing
- itchy nose or throat
- coughs
- cough due to minor throat and bronchial irritation

### Warnings

**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4 doses (120 mL) in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

.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

- to make a child sleepy
- 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.
- with any other drug containing acetaminophen (prescription or non-prescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist
- if you are allergic to acetaminophen or any of the inactive ingredients in this product

## Ask a doctor before use if you have

- liver disease
- glaucoma
- breathing problems
- chronic bronchitis
- a sodium-restricted diet
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough such as occurs with smoking, asthma or emphysema or if cough is accompanied by excessive phlegm (mucus)

#### Ask a doctor or pharmacist before use if you are taking

- sedatives or tranquilizers
- the blood thinning drug warfarin

### When using this product

- do not use more than directed (see overdose warning)
- excitability may occur, especially in children
- marked drowsiness may occur
- avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery
- alcohol, sedatives and tranquilizers may increase drowsiness

## Stop use and ask a doctor if

- redness or swelling is present
- new symptoms occur
- fever gets worse or lasts more than 7 days
- 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**: Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center

(1-800-222-1222) immediately. 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 recommended (see overdose warning)
- do not exceed 4 doses per 24 hours
- use dosage cup provided
- mL= milliliter

age	dose
adults and children 12 yrs and over	30 mL every 6 hrs
children 4 to 11 yrs	do not use unless directed by a doctor
children under 4 yrs	do not use

 When using other Day Time or Night Time products, carefully read each label to ensure correct dosing

#### Other information

- each 15 mL contains: sodium 18 mg
- store at room temperature

#### Inactive ingredients

alcohol, citric acid, FD&C blue 1, FD&C red 40, flavor, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sodium citrate

#### Questions or comments?

call 1-877-753-3935 Monday-Friday 9AM-5PM EST

## **Principal Display Panel**

**COMPARE** TO THE ACTIVE INGREDIENTS **OF VICKS**® **NYQUIL**®\*

## **Nighttime**

#### Cold & Flu

Multi-Symptom Relief

Acetaminophen-Pain Reliever/Fever Reducer

Dextromethorphan HBr-Cough Suppressant)

Doxylamine succinate-Antihistamine

FL OZ (mL)

Cherry Flavor

\*This product is not manufactured or distributed by Procter & Gamble, owner of the registered trademark Vicks® and NyQuil®.

Failure to follow these warnings could result in serius consequences.

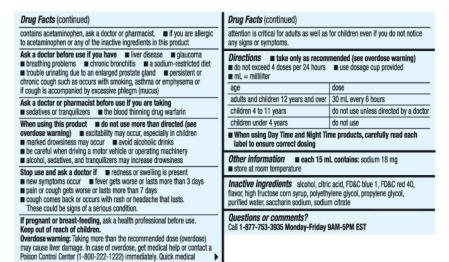
## TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND DOSAGE CUP OR UNDER CAPIS BROKEN OR MISSING.

**DISTRIBUTED BY:** 

FAMILY DOLLAR SERVICES, INC., 10401 MONROE RD, MATTHEWS, NC 28105 USA VISIT FAMILYDOLLAR.COM

#### **Product Label**





## **COLD AND FLU NIGHTTIME**

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55319-342
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	500 mg in 15 mL	
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg in 15 mL	
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg in 15 mL	

Inactive Ingredients			
Ingredient Name	Strength		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
ALCOHOL (UNII: 3K9958V90M)			
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)			
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:55319- 342-10	296 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/29/2016	02/28/2025

Marketing Information				
	Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC monograph final	part341	02/29/2016	02/28/2025

## Labeler - Family Dollar (FAMILY WELLNESS) (024472631)

Revised: 10/2022 Family Dollar (FAMILY WELLNESS)