COLD AND FLU RELIEF MULTI SYMPTOM NIGHTTIME- acetaminophen, dextromethorphan hydrobromide, doxylamine succinate liquid WinCo Foods, LLC

Drug Facts

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 these common cold/flu symptoms:
 - minor aches and pains
 - headache
 - fever
 - sore throat
 - runny nose and sneezing
 - 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 4,000 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 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 is followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

Do not use

- with 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.

Ask a doctor before use if you have

- liver disease
- glaucoma
- cough that occurs with to much phlegm (mucus)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis, or emphysema
- trouble urinating due to an enlarged prostate gland

Ask a doctor or pharmacist before use if you are taking

- sedatives or tranquilizers
- the blood thinning drug warfarin

When using this product

- 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

- pain or cough gets worse or lasts more than 7 days
- new symptoms occur
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- 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) right away. Quick 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 (see Overdose warning)
- do not take more than 4 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device.
- mL= milliliter
- keep dosing cup with product
 - o adults and children 12 years and over: 30 mL every 6 hours
 - children under 12 years of age: do not use
- When using other Day Time or Night Time products, carefully read each label to insure correct dosing

Other information

- each 30 mL contains: potassium 5 mg
- each 30 mL contain: sodium 19 mg
- store between 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

acesulfame potassium, alcohol, citric acid, D&C yellow #10, FD&C green #3, FD&C yellow #6, flavor, high fructose corn syrup, polyethylene glycol, propylene glycol, purified water, saccharin sodium, sodium citrate

Questions or comments?

Call 800-824-1706 Monday-Friday 9am-4pm MST

Principal Display Panel

*Compare to the active ingredients in Vicks® NyQuil® Cold & Flu

Multi-Symptom

Cold & Flu Relief

NIghttime

Acetaminophen/Pain Reliever-Fever Reducer

Dextromethorphan HBr/Cough Suppressant

Doxylamine succinate/Antihistamine

For ages 12 and over

Alcohol 10%

Original Flavor

FL OZ (mL)

*This product is not manufactured or distributed by THE PROCTER & GAMBLE

COMPANY.VICKS® and NYQUIL® are registered trademarks of THE PROCTER & GAMBLE COMPANY.

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

DISTRIBUTED BY: WINCO FOODS, LLC, BOISE, ID 83704

Product Label



WINCO FOODS Nighttime Multi-Symptom Cold & Flu Relief

COLD AND FLU RELIEF MULTI SYMPTOM NIGHTTIME

acetaminophen, dextromethorphan hydrobromide, doxylamine succinate liquid

Product Information	duct Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67091-258	
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)	ACETAMINOPHEN	650 mg in 30 mL		
DEXTROMETHORPHAN HYDROBROMIDE (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		
UIVII.93QD//JKFL/		III 30 IIIL		

Inactive Ingredients		
Ingredient Name	Strength	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
ACESULFAME POTASSIUM (UNII: 230V73Q5G9)		
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)		
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)		
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)		
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC:67091- 258-54	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/31/2014	
NDC:67091- 258-23	237 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/31/2014	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M012	12/31/2014		

Labeler - WinCo Foods, LLC (056098817)

Revised: 5/2024 WinCo Foods, LLC