TUSSIN CF NON DROWSY MULTI SYMPTOM- dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid WinCo Foods, LLC

Drug Facts

Active ingredients (in each 10 mL)

Dextromethorphan HBr 20 mg Guaifenesin 200 mg Phenylephrine HCl 10 mg

Purposes

Cough suppressant

Expectorant

Nasal decongestant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes
- temporarily relieves these symptoms occurring with a cold:
 - nasal congestion
 - o cough due to minor throat and bronchial irritation

Warnings

Do not use

• 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

- heart disease
- diabetes
- high blood pressure
- thyroid disease
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema
- cough that occurs with too much phlegm (mucus)
- trouble urinating due to enlarged prostate gland

Ask a doctor or pharmacist before use if you are

taking any other oral nasal decongestant or stimulant.

When using this product,

do not use more than directed.

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- symptoms do not get better within 7 days or are accompanied by fever
- cough lasts more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache.

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 (1800-222-122) right away.

Directions

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device.
- keep dosing cup with product
- mL = milliliter
- this adults product is not intended for use in children under 12 years of age
- adult and children 12 years and over: 10 mL every 4 hours
- children under 12 years: do not use

Other information

• store between 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

anhydrous citric acid, FD&C red #40, flavor, glycerin, lactic acid, menthol, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose

Questions?

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

Principal Display Panel

*Compare to the active ingredients in Robitussin® Peak Cold Multi-Symptom Cold CF

NON-DROWSY

Tussin CF

Multi-Symptom

DEXTROMETHORPHAN HBr / COUGH SUPPRESSANT

GUAIFENESIN / EXPECTORANT

PHENYLEPHRINE HCI / NASAL DECONGESTANT

Relieves: Cough, Stuffy Nose, Chest Congestion & Mucus

Adult Formula, For Ages 12 & Over

Alcohol Free

FL OZ (mL)

*This product is not manufactured or distributed by Pfizer Consumer Healthcare, distributors of Robitussin® Peak Cold Multi-Symptom Cold CF.

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

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

www.wincofoods.com

Package Label



TUSSIN CF NON DROWSY MULTI SYMPTOM

dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67091-266
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 10 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 10 mL

PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE -	PHENYLEPHRINE	10 mg
UNII:1WS297W6MV)	HYDROCHLORIDE	in 10 mL

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
LACTIC ACID (UNII: 33X04XA5AT)		
MENTHOL (UNII: L7T10EIP3A)		
SORBITOL (UNII: 506T60A25R)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67091- 266-18	1 in 1 BOX	12/31/2014	
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

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