TUSSIN COUGH DM SUGAR FREE- dextromethorphan hbr, guaifenesin liquid QUALITY CHOICE (Chain Drug Marketing Association)

Drug Facts

Active ingredients (in each 10 mL)

Dextromethorphan HBr,20 mg Guaifenesin,200 mg

Purposes

Cough suppressant

Expectorant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

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

- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis, or emphysema

Stop use and ask a doctor if

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 (1-800-222-1222) 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
- adults 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°). Do not refrigerate.

Inactive ingredients

acesulfame potassium, citric acid, flavors, glycerin, methylparaben, polyethylene glycol, povidone, propylene glycol, purified water, saccharin sodium, sodium benzoate

Questions or comments?

Call **1-248-449-9300** Monday-Friday 9AM-5PM EST

Principal Display Panel

*Compare to the Active Ingredients in Robitussin® Peak Cold Sugar-Free Cough + Chest Congestion DM

Sugar-Free

Non-Drowsy

Adult

TUSSIN DM

Cough & Chest Congestion

Dextromethorphan HBr 20 mg Cough Suppressant

Guaifenesin 200 mg Expectorant

Relieves:

Cough & Mucus

For Ages 12 Years & Over

Alcohol-Free

DM Sugar-Free Formula Specially Formulated for Diabetic Use

FL OZ (mL)

Dosing Cup Included

*This product is not manufactured or distributed by Pfizer Consumer Healthcare, distributors of Robitussin® Peak Cold Sugar-Free Cough+Chest Congestion DM.

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

Distributed by C.D.M.A., Inc.© 43157 W 9 Mile Rd Novi, MI 48375 www.qualitychoice.com

Package Label





Purnoses

TUSSIN COUGH DM SUGAR FREE

dextromethorphan hbr, guaifenesin liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63868-869

Route of Administration ORAL

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

Inactive Ingredients					
Ingredient Name	Strength				
ACESULFAME POTASSIUM (UNII: 230V73Q5G9)					
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)					
GLYCERIN (UNII: PDC6A3C0OX)					
METHYLPARABEN (UNII: A2I8C7HI9T)					
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)					
POVIDONE (UNII: FZ 989GH94E)					
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)					
WATER (UNII: 059QF0KO0R)					
SACCHARIN SODIUM (UNII: SB8ZUX40TY)					
SODIUM BENZOATE (UNII: OJ245FE5EU)					

Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
	NDC:63868- 869-04	1 in 1 BOX	08/31/2018				
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product					

Marketing In	formation		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	08/31/2018	