

**FAMILY CARE COUGH AND COLD- dextromethorphan hydrobromide, and guaifenesin liquid
United Exchange Corp.**

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.

Active ingredients (in each 10 mL)	Purpose
Dextromethorphan HBr, USP 20 mg.....	Cough suppressant
Guaifenesin, USP 200 mg.....	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 brnchial 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 until 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)
- or a 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 right away.

Directions

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided. Keep dosing cup with product
- mL = milliliter
- this adult product is not intended for use in children under 12 years of age

age	dose
adults and children 12 years and over	10 mL every 4 hours
children under 12 years	do not use

Other information

- each 10 mL contains: sodium 3 mg
- Store at 20° to 25°C (68° to 77°F). Do not refrigerate.

Inactive ingredients

citric acid hydrate, FD&C red no. 40, fruit mixing essence, glycerin, high fructose corn syrup, L-

menthol, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate hydrate, sucralose

Distributed by:

United Exchange Corp.

17211 Valley View Ave.

Cerritos, CA 90703 USA

Made in Korea

TAMPER EVIDENT: Do not use if printed shrinkband is missing or broken. Failure to follow these warnings could result in serious consequences.

Drug Facts

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Compare to the active ingredients in ROBITUSSIN® Cough + Chest Congestion DM

FAMILY CARE

COUGH & COLD

Relieves: ■ Cough ■ Mucus

ADULT DM

NON-DROWSY • ALCOHOL FREE

PARENTS: Should be 18 or older to purchase. Learn about teen medicine abuse. www.StopMedicineAbuse.org

DEXTROMETHORPHAN HBR
GUAIFENESIN

4 FL OZ (118 mL)

Drug Facts (continued)

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*This product is not manufactured or distributed by Pfizer, owner of the registered trademark Robitussin® Cough + Chest Congestion DM.

Distributed by:
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FAMILY CARE COUGH AND COLD

dextromethorphan hydrobromide, and guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-633
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	40 mg in 10 mL

GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)

GUAIFENESIN

400 mg
in 10 mL**Inactive Ingredients**

Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
MENTHOL (UNII: L7T10EIP3A)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-633-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	07/29/2015	

Labeler - United Exchange Corp. (840130579)

Revised: 7/2015

United Exchange Corp.