

**FAMILY CARE MULTI SYMPTOM COLD- dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride 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.*

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| Active ingredient (in each 10 mL)    | Purpose            |
|--------------------------------------|--------------------|
| Gextromethorphan HBr, USP 20 mg..... | Cough suppressant  |
| Guaifenesin, USP 200 mg.....         | Expectorant        |
| Phenylephrine HCl, USP 10 mg.....    | 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
- 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 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 or pharmacist before us 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

- you get nervous, dizzy, or sleepless
  - 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 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

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

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**Other information**

- each 10 mL contains: sodium 2 mg

- store at 20° to 25°C (68° to 77°F)
- Do not refrigerate

Inactive ingredients

citric acid hydrate, D-sorbitol, FD&C red no. 40, glycerin, high fructose corn syrup, lemon essence, L-menthol, propylene glycol, purified water, sodium benzoate, sodium citrate hydrate, sucralose

Distributed by:

United Exchange Corp.

17211 Valley View Ave.

Cerritos, CA 90703 U.S.A

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.

**Compare to the active ingredients in ROBITUSSIN® Multi-Symptom CF**

**FAMILY CARE**  
**MULTI-SYMP TOM**  
**COLD**  
Relieves: ■ Nasal Congestion ■ Cough ■ Mucus  
**ADULT** **CF**

**NON-DROWSY**  
Should be 18 or older to purchase  
**PARENTS:**  
Learn about teen medicine abuse  
[www.StopMedicineAbuse.org](http://www.StopMedicineAbuse.org)  
For Ages 12 & Over 4 FL OZ (118 mL)  
DEXTROMETHORPHAN HBr  
GUAIFENESIN  
PHENYLEPHRINE HCl

**Drug Facts (continued)**  
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.  
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\*This product is not manufactured or distributed by Pfizer, owner of the registered trademark Robitussin® Multi-Symptom CF.  
Distributed by: United Exchange Corp. 17211 Valley View Ave. Cerritos, CA 90703 USA www.uccorp.com Toll Free: 1 800 814 8028 Made in Korea

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**FAMILY CARE MULTI SYMPTOM COLD**

dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride liquid

**Product Information**

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:65923-632 |
| <b>Route of Administration</b> | ORAL           |                           |               |

**Active Ingredient/Active Moiety**

| Ingredient Name   | Basis of Strength                | Strength          |
|---|----------------------------------|-------------------|
| <b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH)<br>(DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN<br>HYDROBROMIDE | 10 mg<br>in 5 mL  |
| <b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)                           | GUAIFENESIN                      | 100 mg<br>in 5 mL |
| <b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)         | PHENYLEPHRINE<br>HYDROCHLORIDE   | 5 mg<br>in 5 mL   |

**Inactive Ingredients**

| Ingredient Name                                    | Strength |
|--|----------|
| <b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)  |          |
| <b>SORBITOL</b> (UNII: 506T60A25R)                 |          |
| <b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)      |          |
| <b>GLYCERIN</b> (UNII: PDC6A3C0OX)                 |          |
| <b>HIGH FRUCTOSE CORN SYRUP</b> (UNII: XY6UN3QB6S) |          |
| <b>MENTHOL</b> (UNII: L7T10EIP3A)                  |          |
| <b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)         |          |
| <b>WATER</b> (UNII: 059QF0K00R)                    |          |
| <b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)          |          |
| <b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)                |          |

**Packaging**

| # | Item Code        | Package Description  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:65923-632-04 | 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 final | part341                                  | 07/31/2015           |                    |

**Labeler** - United Exchange Corp. (840130579)

Revised: 7/2015

United Exchange Corp.