

**CHILDRENS ROBITUSSIN COUGH AND CHEST CONGESTION DM-  
dextromethorphan hydrobromide, guaifenesin liquid  
Haleon US Holdings LLC**

-----  
**Drug Facts**

**Active ingredients (in each 5 ml)**

Dextromethorphan HBr, USP 5 mg

Guaifenesin, USP 100 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 right away.

**Directions**

- measure only with dosing cup provided
  - keep dosing cup with product
  - ml = milliliter
  - do not take more than 6 doses in any 24-hour period
-

<b>age</b>	<b>dose</b>
children under 4 years	do not use
children 4 to under 6 years	5 ml every 4 hours
children 6 to under 12 years	10 ml every 4 hours
adults and children 12 years and over	20 ml every 4 hours

### ***Other information***

- **each 5 ml contains:**sodium 3 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

### ***Inactive ingredients***

anhydrous citric acid, artificial flavor, FD&C blue no. 1, FD&C red no. 40, glycerin, natural flavor, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose

### ***Questions or comments?***

call weekdays from 9 AM to 5 PM EST at **1-800-762-4675**

For most recent product information, **visit [www.robitussin.com](http://www.robitussin.com)**

Distributed by:

Pfizer, Madison, NJ 07940 USA

## **PRINCIPAL DISPLAY PANEL**

***NEW***

***Dosing***

***Information***

**Children's**

**Robitussin<sup>®</sup>**

**NOW FOR AGES 4 & OVER**

**Cough & Chest**

**Congestion**

**DM**

**DEXTROMETHORPHAN HBr**

**(Cough Suppressant)**

**GUAIFENESIN (Expectorant)**

**Relieves:**

✓ **Chest Congestion/Mucus**

✓ **Cough**

**Non-Drowsy**

**grape**

flavor

4 FL OZ

(118 ml)



## CHILDRENS ROBITUSSIN COUGH AND CHEST CONGESTION DM

dextromethorphan hydrobromide, guaifenesin liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0031-8715
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	5 mg in 5 mL
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>PROPYLENE GLYCOL, (R)-</b> (UNII: 602HN5L69H)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CITRATE, UNSPECIFIED FORM</b> (UNII: 1Q73Q2JULR)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	

### Product Characteristics

<b>Color</b>	purple (purple)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	GRAPE (grape)	<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0031-8715-10	1 in 1 CARTON	07/01/2014	
1		118 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/01/2014	

**Labeler** - Haleon US Holdings LLC (079944263)