

**CHILDRENS ROBITUSSIN ELDERBERRY COUGH AND CHEST CONGESTION DM-
dextromethorphan hbr, guaifenesin solution
Haleon US Holdings LLC**

Drug Facts

Active ingredients (in each 20 mL)

Dextromethorphan HBr 20 mg

Guaifenesin 400 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

- 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

age	dose
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 20 mL contains:**potassium 5 mg, sodium 21 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

anhydrous citric acid, black elderberry juice concentrate (for flavor), carboxymethylcellulose sodium, glycerin, liquid glucose, maltodextrin, natural and artificial flavors, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate, sodium gluconate, sucralose, xanthan gum, zinc gluconate

Questions or comments?

call weekdays from 8 AM to 6 PM EST at **1-800-245-1040**

Additional information

Distributed by: Haleon, Warren, NJ 07059

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For most recent product information, visit www.robitussin.com

Pat. Info www.productpats.com

Made in Canada

PARENTS:

Learn about teen medicine abuse

www.StopMedicineAbuse.org

Packaged with Tamper-Evident bottle cap.

Do Not Use if breakable ring is separated or missing.

PRINCIPAL DISPLAY PANEL

Children’s

Robitussin

Elderberry

**Cough & Chest
Congestion DM**

DEXTROMETHORPHAN HBr (Cough Suppressant)

GUAIFENESIN (Expectorant)

- Controls Cough
- Relieves Chest Congestion
- Thins & Loosens Mucus

Taste the

Real Elderberry

4 FL OZ (118 mL)

For Ages 4+

62000000206394 Front Carton

Children's

Robitussin

Elderberry



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0031-2098
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 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SODIUM GLUCONATE (UNII: R6Q3791S76)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	
ZINC GLUCONATE (UNII: U6WSN5SQ1Z)	

Product Characteristics

Color	purple (dark-purple to dark purple-brown)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0031-2098-01	1 in 1 CARTON	06/14/2021	
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	06/14/2021	

Labeler - Haleon US Holdings LLC (079944263)