

**CHILDRENS ROBITUSSIN HONEY NIGHTTIME COUGH DM- dextromethorphan
hbr, doxylamine succinate solution**
Haleon US Holdings LLC

Drug Facts

Active ingredients (in each 10mL)

Dextromethorphan HBr, USP 15 mg

Doxylamine Succinate, USP 6.25 mg

Purposes

Cough Suppressant

Antihistamine

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
 - runny nose
 - sneezing
 - itchy, watery eyes
 - itching of the nose or throat
- controls the impulse to cough to help you sleep

Warnings

Do not use

- to sedate a child or to make a child sleepy
- 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

- trouble urinating due to an enlarged prostate gland
- glaucoma
- a cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis or emphysema

Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers

When using this product

- **do not use more than directed**

- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

Stop use and ask a doctor ifcough 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 4 doses in any 24-hour period

age	dose
children under 6 years	do not use
children 6 to under 12 years	10 mL every 6 hours
adults and children 12 years and older	20 mL every 6 hours

Other information

- **each 10 mL contains:** sodium 10 mg
- store at 20–25°C (68–77°F)

Inactive ingredients

anhydrous citric acid, blueberry juice concentrate, carboxymethylcellulose sodium, glycerin, lactic acid, natural and artificial flavors, natural grade A honey, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate, sodium gluconate, sucralose, xanthan gum, zinc gluconate

Questions or comments?

call weekdays from 9 AM to 5 PM EST at **1-800-245-1040**

Additional Information

Packaged with Tamper-Evident bottle cap. Do Not Use if breakable ring is separated or missing.

Dist. by: Haleon, Warren, NJ 07059

©2024 Haleon or licensor.

Trademarks owned or licensed by Haleon.

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

Pat. Info www.productpats.com

Made in Canada

PRINCIPAL DISPLAY PANEL

**Children's
Robitussin
Honey
For Ages 6+**

**Nighttime
Cough
DM**

**DEXTROMETHORPHAN HBR
(COUGH SUPPRESSANT)
DOXYLAMINE SUCCINATE (ANTIHISTAMINE)**

ALCOHOL
FREE

LONG-ACTING

- Relieves:
- Cough up to **8**hours
- Runny nose

Taste the
Real Honey

TRUE
SOURCE
CERTIFIED
HONEY ✓

4 FL OZ (118 mL)



CHILDRENS ROBITUSSIN HONEY NIGHTTIME COUGH DM

dextromethorphan hbr, doxylamine succinate solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0031-8762
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII: 7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg in 10 mL
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII: 95QB77JKPL)	DOXYLAMINE SUCCINATE	6.25 mg in 10 mL

Inactive Ingredients				
Ingredient Name			Strength	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)				
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)				
GLYCERIN (UNII: PDC6A3C0OX)				
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)				
HONEY (UNII: Y9H1V576FH)				
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		Score		
Shape		Size		
Flavor	BERRY	Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0031-8762-12	1 in 1 CARTON	05/27/2019	
1		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 Drug		M012	05/27/2019	

Labeler - Haleon US Holdings LLC (079944263)