CHILDRENS ROBITUSSIN NIGHTTIME COUGH LONG-ACTING DM-
chlorpheniramine maleate, dextromethorphan hydrobromide solution
Haleon US Holdings LLC

Drug Facts

Active ingredients (in each 10 mL)

Chlorpheniramine maleate, USP 2 mg

Dextromethorphan HBr, USP 15 mg

Purposes

Antihistamine

Cough suppressant

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

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 aretaking 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 6 mg
- store at 20-25°C (68-77°F)

Inactive ingredients

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

Questions or comments?

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

Additional Information

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

Children’s Robitussin liquid is specially formulated to provide soothing action, control your child’s cough plus relieve other cold symptoms.

Should be 18 or older to purchase

PARENTS:

Learn about teen medicine abuse

www.StopMedicineAbuse.org

Use dosage cup included

Distributed by: GSK Consumer Healthcare,

Warren, NJ 07059

For most recent product information,

visit www.robitussin.com

Made in Canada

Principal Display Panel

Children's

Robitussin

AGES 6 & OVER

Nighttime
Cough

Long-Acting

DM

CHLORPHENIRAMINE MALEATE (Antihistamine)
DEXTROMETHORPHAN HBr (Cough Suppressant)

Relieves Cough up to 8 Hours
Runny Nose

Alcohol-Free

Fruit punchflavor

4 FL OZ (118 mL)

PAA172103 Front Carton

CHILDRENS ROBITUSSIN NIGHTTIME COUGH LONG-ACTING DM

chlorpheniramine maleate, dextromethorphan hydrobromide solution

Product Information			

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0031-8692
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg in 10 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
LACTIC ACID, UNSPECIFIED FORM (UNII: 33X04XA5AT)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color	red (red)	**Score**	
Shape		**Size**	
Flavor	FRUIT PUNCH	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0031-8692-13	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)