CHILDRENS ROBITUSSIN COUGH AND COLD CF- dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride liquid
GlaxoSmithKline Consumer Healthcare Holdings (US) LLC

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.

Drug Facts

Active ingredients (in each 10 mL)

Dextromethorphan HBr, USP 10 mg

Guaifenesin, USP 100 mg

Phenylephrine HCl, USP 5 mg

Purposes

Cough suppressant

Expectorant

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 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

1. heart disease
2. high blood pressure
3. thyroid disease
4. diabetes
5. trouble urinating due to an enlarged prostate gland
6. cough that occurs with too much phlegm (mucus)

7. cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema

Ask a doctor or pharmacist before use 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

- 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

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 10 mL contains:** sodium 7 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

anhydrous citric acid, artificial flavor, FD&C red no. 40, glycerin, 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

Children’s Robitussin liquid is especially 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

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

FOR AGES 4 & OVER

Cough & Cold

CF

DEXTROMETHORPHAN HBr
(Cough Suppressant)
GUAIFENESIN (Expectorant)
PHENYLEPHRINE HCl (Nasal Decongestant)

Relieves:

- **Cough**
- **Chest Congestion/Mucus**
- **Stuffy Nose**

Non-Drowsy

grape
flavor

4 FL OZ
(118 mL)

PAA172104 Front Carton

CHILDRENS ROBITUSSIN COUGH AND COLD CF

dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride liquid

Product Information			

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0031-8716
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	5 mg in 5 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	50 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	2.5 mg in 5 mL

Inactive Ingredients

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

Product Characteristics

Color	RED (clear red)	**Score**	
Shape		**Size**	
Flavor	GRAPE	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0031-8716-12	1 in 1 CARTON	09/01/2009	
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 final	part341	09/01/2009	

Labeler - GlaxoSmithKline Consumer Healthcare Holdings (US) LLC (079944263)

Revised: 7/2023 GlaxoSmithKline Consumer Healthcare Holdings (US) LLC