ROBITUSSIN CF- dextromethorphan hydrobromide, guaifenesin, pseudoephedrine hydrochloride liquid GlaxoSmithKline Consumer Healthcare Holdings (US) LLC

Robitussin® CF

Active ingredients (in each 5 mL tsp)	Purpose
Dextromethorphan HBr, USP 10 mg	Cough suppressant
Guaifenesin, USP 100 mg	Expectorant
Pseudoephedrine HCl, USP 30 mg	Nasal decongestant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive
- 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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema

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.

Directions

do not take more than 4 doses in any 24-hour period

age	dose
adults and children	2 teaspoonfuls every 4
12 years and over	hours
children 6 years to	1 teaspoonful every 4
under 12 years	hours
under 6 years	Do not use

Inactive ingredients

citric acid, FD&C red no. 40, flavors, glycerin, propylene glycol, saccharin sodium, sodium benzoate, sorbitol, water.

Keep out of reach of children. In case of overdose, get medical help right away.

Other information

- store at 20-25°C (68-77°F)
- dosage cup provided

MADE IN U.S.A.

PRINCIPAL DISPLAY PANEL - 118 mL Bottle Carton

Wyeth[®]

Robitussin®

CF

NASAL DECONGESTANT COUGH SUPPRESSANT EXPECTORANT

- CLEARS NASAL STUFFINESS
- CONTROLS COUGHS
- LOOSENS and RELIEVES CHEST CONGESTION

COUGH FORMULA For Children and Adults

4 FL OZ (118 mL)

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Robitussin

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Wyeth®

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NEW FORMULA

Robitussin

NASAL DECONGESTANT COUGHSUPPRESSANT EXPECTORANT

CHOOSE the correct Robitussin Formula for your Symptoms.

Chest Congestion with Dry Cough?

 Loosens Chest Congestion to Help Coughs Clear Chest.

Frequent Coughs + Chest Congestion?

Robitussin-DM

- Controls and Suppresses Coughing.
- Loosens and Relieves Chest Congestion.

Nasal Stuffiness + Chest Congestion with **Dry Cough?**

Robitussin-PE

- Clears Nasal Passages to Restore Freer Breathing Through the Nose.
- Loosens and Relieves Chest Congestion.

Nasal Stuffiness + Frequent Coughs + Chest Congestion?

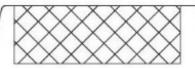
- Clears Nasal Passages to Restore Freer Breathing Through the Nose. Controls and Suppresses Coughing.
- · Relieves Chest Congestion.
- · Sugar-Free.

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MADE IN U.S.A.

LOT: EXP:

MFD:



EVIDENT BOTTLE CAP. PA CKA GED WITH TAMPER-

NEW FORMULA

Robitussi

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NEW FORMULA

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NASAL DECONGESTANT COUGH SUPPRESSANT EXPECTORANT

Why Are Robitussin Products Chosen More Often?



RECOMMENDED by more Doctors and Pharmacists than any other cough formula.



RELIABLE and EFFECTIVE formulas for the WHOLE FAMILY.

NON-DROWSY and NON-NARCOTIC formulas.

Richmond Division of Wyeth 2248-2300 Darbytown Rd. Richmond, VA 23231

Keep out of reach of children. In case of overdose, get medical help right away.

Other information

- store at 20-25°C (68-77°F) يغزن في درجة حرارة الغرقة بين ٢٠ إلى ٢٥ درجة مئوية
- · dosage cup provided



PAA019124

ROBITUSSIN CF

dextromethorphan hydrobromide, guaifenesin, pseudoephedrine hydrochloride liquid

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0031-8678	
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL		
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL		
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	30 mg in 5 mL		

Inactive Ingredients				
Ingredient Name	Strength			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
FD&C RED NO. 40 (UNII: WZB9127XOA)				
GLYCERIN (UNII: PDC6A3C0OX)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				
SORBITOL (UNII: 506T60A25R)				
WATER (UNII: 059QF0KO0R)				

F	Packaging					
#	tem Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:0031-8678- 82	1 in 1 CARTON	07/01/2015			
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product				

Marketing Information					
Marketing Application Number or Monograph Marketing Start Marketing End Category Citation Date Date					
Export only		07/01/2015			

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

Revised: 1/2023 GlaxoSmithKline Consumer Healthcare Holdings (US) LLC