

**ROBITUSSIN PEAK COLD MULTI-SYMPTOM COLD- 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 20 mg

Guaifenesin, USP 200 mg

Phenylephrine HCl, USP 10 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**

- 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

**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***

- 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
- this adult product is not intended for use in children under 12 years of age

<b>age</b>	<b>dose</b>
adults and children 12 years and over	10 ml every 4 hours
children under 12 years	do not use

***Other information***

- **each 10 ml contains:** sodium 6 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

***Inactive ingredients***

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

***Questions or comments?***

call weekdays from 9 AM to 5 PM EST at **1-800-762-4675**

For most recent product information,

**visit [www.robitussin.com](http://www.robitussin.com)**

Distributed by:  
Pfizer, Madison, NJ 07940 USA

**PRINCIPAL DISPLAY PANEL**

**ADULT**

**Robitussin®**

**PEAK COLD**

**Multi-Symptom  
Cold**

**DEXTROMETHORPHAN HBr (Cough Suppressant)**

**GUAIFENESIN (Expectorant)**

**PHENYLEPHRINE HCl (Nasal Decongestant)**

**Relieves:**

- **Cough**
- **Nasal Congestion**
- **Mucus**

**Non-Drowsy CF**

For Ages 12 & Over

4 FL OZ  
(118 ml)



## ROBITUSSIN PEAK COLD MULTI-SYMPATOM COLD

dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0031-8742
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 5 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

<b>MENTHOL, UNSPECIFIED FORM</b> (UNII: L7T10EIP3A)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CITRATE, UNSPECIFIED FORM</b> (UNII: 1Q73Q2JULR)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Product Characteristics

<b>Color</b>	RED (clear red liquid)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	BERRY (berry-citrus flavor) , CITRUS	<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0031-8742-14	1 in 1 CARTON	06/12/2011	
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0031-8742-20	1 in 1 CARTON	06/12/2011	
2		237 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 FINAL	part341	06/12/2011	

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

Revised: 12/2022

GlaxoSmithKline Consumer Healthcare Holdings (US) LLC