TUSSIN CF - dextromethorphan hbr guaifenesin phenylephrine hcl solution
Strategic Sourcing Services

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.

Tussin CF
Dextromethorphan HBr Guaifenesin Phenylephrine HCl

Active ingredients (in each 10 mL)

Dextromethorphan HBrDextromethorphan 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 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 (1-800-222-1222).

Directions

■ mL = milliliter
■ do not take more than 6 doses in any 24 hour period
■ this adult strength product is not intended for use in children under 12 years of age
■ measure only with dosing cup provided
■ keep dosing cup with product

AGE	DOSE
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

■ Keep carton for full Direction for use

Inactive ingredients

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

Questions or comments?

Call **833-358-6431**Monday to Friday 9:00am to 7:00pm EST

Distributed by: McKesson Corp., via Strategic Sourcing Services LLC.
Memphis, TN 38141
Money Back Guarantee
www.fosterandthrive.com

*This product is not manufactured or distributed by Pfizer,
owner of the registered trademark Robitussin ®

PRINCIPAL DISPLAY PANEL

Dextromethorphan HBr Guaifenesin Phenylephrone HCL-237 mL

TUSSIN CF

dextromethorphan hbr guaifenesin phenylephrine hcl solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70677-1187
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 10 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 10 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL (UNII: L7T10EIP3A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70677-1187-1	1 in 1 CARTON	08/09/2023	
1		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	08/09/2023	

Labeler - Strategic Sourcing Services (116956644)

Establishment

Name	Address	ID/FEI	Business Operations
AptaPharma Inc.		790523323	manufacture(70677-1187)

Revised: 8/2023 Strategic Sourcing Services