TUSSIN MULTI SYMPTOM COLD CF- dextromethorphan hbr guaifenesin phenylephrine hcl liquid
QUALITY CHOICE (Chain Drug Marketing Association)

Drug Facts

Active ingredients (in each 10 mL)

Dextromethorphan HBr 20 mg

Guaifenesin 200 mg

Phenylephrine HCl 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
- diabetes
- thyroid
- 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

- nervousness, dizziness, or sleeplessness occur
- 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 (1800-222-1222) right away.

Directions

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device.
- keep dosing cup with product
- mL = milliliter
- this adult product is not intended for use in children under 12 years of age
- adult and children 12 years and over: 10 mL every 4 hours
- children under 12 years: do not use

Other information

- store between 20-25°C (68°-77°F). Do not refrigerate.

Inactive ingredients

anhydrous citric acid, FD&C red #40, flavor, glycerin, lactic acid, menthol, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose

Questions or comments?

Call **1-800-935-2362** Monday-Friday 9AM-5PM EST

Principal display panel

*Compare to the Active Ingredients in Robitussin® Peak Cold Multi-Symptom Cold CF

Adult

Tussin CF

Multi-Symptom Cold

Dextromethorphan HBr

Cough suppressant

Guaifenesin

Expectorant

Phenylephrine HCl

Nasal decongestant

Relieves:

- Cough
- Mucus
- Nasal congestion

For ages 12 years & over

Alcohol Free

Non-Drowsy

FL OZ (mL)

Dosing Cup Included

*This product is not manufactured or distributed by Haleon group of companies, distributors of Robitussin® Peak Cold Multi-Symptom Cold CF.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF PRINTED SAFETY SEAL AROUND BOTTLE OR UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Distributed by CDMA, Inc.

Novi, MI 48375

Package Label

QUALITY CHOICE Adult Multi-Symptom Cold

TUSSIN MULTI SYMPTOM COLD CF

dextromethorphan hbr guaifenesin phenylephrine hcl liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-155
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)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
LACTIC ACID (UNII: 33X04XA5AT)	
MENTHOL (UNII: L7T10EIP3A)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-155-04	1 in 1 BOX	07/31/2024	
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/31/2024	

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Revised: 5/2024 QUALITY CHOICE (Chain Drug Marketing Association)