TUSSIN DM COUGH SUPPRESSANT/EXPECTORANT- dextromethorphan hydrobromide, guaifenesin liquid Chain Drug Consortium, LLC

Active ingredients

Drug Facts

Active ingredients (in each 5 mL tsp)

Dextromethorphan HBr, USP 10 mg Guaifenesin, USP 100 mg

Purpose

Cough Suppressant Expectorant

Keep out of reach of children

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away.

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

Warnings

Do not use

- in a child under 12 years of age
- 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

- cough that occurs with too much phlegm mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema

Stop use and ask a doctor if

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 6 doses in any 24-hour period

Other information

- Keep carton for full Directions for use.
- store at 20-25 ° C (68-77 ° F)
- do not refrigerate
- dosage cup provided
- sodium 3 mg per teaspoonful

Inactive ingredients

anhydrous citric acid, dextrose, FD and C red no.40, flavor, glycerin, high fructose corn syrup, menthol, purified water, saccharin sodium, sodium benzoate

Questions?

Call weekdays from 9:30 AM to 4:30 PM EST at

1-877-798-5944

Product Label

NDC 68016-018-04

*COMPARE TO THE ACTIVE INGREDIENTS IN ROBITUSSIN® PEAK COLD COUGH and CHEST CONGESTION DM

Premier Value® Tussin DM

Dextromethorphan HBr Guaifenesin

COUGH SUPPRESSANT /

EXPECTORANT

Helps to Loosen Chest Congestion

NON-DROWSY

Cough Formula for ages 12 and over

4 FL OZ (118 mL)

INDEPENDENTLY TESTED SATISFACTION GUARANTEED PV DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSING

If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refundd.

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

DISTRIBUTED BY: CHAIN DRUG CONSORTIUM 3301 NW BOCA RATON BLVD SUITE 101, BOCA RATON, FL 33431

BX-003

Drug Facts

Active ingredients

Purpose

Expectorant

(in each 5 mL tsp) Dextromethorphan HBr, USP 10 mg....Cough Suppressant Guaifenesin, USP 100 mg.

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

Warnings Do not use

■ in a child under 12 years of age

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

cough that occurs with too much phlegm (mucus) cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema

Stop use and ask a doctor if 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

Age	Dose
adults & children 12 years & over	2 teaspoonfuls every 4 hours
children under 12 years	do not use

Drug Facts (continued)

Other information

- Keep carton for full Direction for use.
- store at 20-25°C (68-77°F)
- do not refrigerate
- dosage cup provided
- sodium 3 mg per teaspoonful

Inactive ingredients

Anhydrous citric acid, dextrose, FD&C red # 40, flavor, glycerin, high fructose com syrup, menthol, purified water, saccharin sodium, sodium benzoate.

Questions?

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BX-003



Tussin DM

Dextromethorphan HBr/ Guaifenesin

COUGH SUPPRESSANT/ EXPECTORANT

Helps Loosen Chest Congestion

NON-DROWSY

Cough Formula for ages 12 & over

4 FL OZ (118 mL)



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TUSSIN DM COUGH SUPPRESSANT/EXPECTORANT

dextromethorphan hydrobromide, quaifenesin liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68016-018

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
(DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL

Inactive Ingredients			
Ingredient Name	Strength		
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
DEXTROSE (UNII: IY9XDZ 35W2)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
GLYCERIN (UNII: PDC6A3C0OX)			
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)			
MENTHOL (UNII: L7T10EIP3A)			
WATER (UNII: 059QF0KO0R)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			

Product Characteristics				
Color		Score		
Shape		Size		
Flavor	CHERRY	Imprint Code		
Contains				

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:68016- 018-04	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/01/2012	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/01/2012	

Labeler - Chain Drug Consortium, LLC (101668460)

Registrant - AptaPharma Inc. (790523323)

Establishment				
Name	Address	ID/FEI	Business Operations	
AptaPharma Inc.		790523323	manufacture(68016-018)	

Revised: 12/2023 Chain Drug Consortium, LLC