

QCH ADULT TUSSIN DM 542 - dextromethorphan hbr, guaifenesin liquid
Chain Drug Marketing Association Inc.

QCH Adult Tussin DM

ACTIVE INGREDIENT(S)

Dextromethorphan HBr, USP 20 mg

Guaifenesin, USP 200 mg

PURPOSE

Cough suppressant

Expectorant

USE(S)

- 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

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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

- 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 DOCTOR IF

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

shake well before using

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

Age	Dose
adults & children 12 years & over	20 mL every 4 hours
children under 12 years	do not use

OTHER INFORMATION

- **each 20 mL contains:** sodium 15 mg
- store at 20-25°C (68-77°F)
- do not refrigerate

INACTIVE INGREDIENTS

carboxymethylcellulose sodium, citric acid, FD&C blue 1 dye, FD&C red 40 dye, flavor, glycerin, high fructose corn syrup, menthol, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate, sucralose, xanthan gum.

PRINCIPAL DISPLAY PANEL

NDC 83324-023-04

QUALITY CHOICE

***Compare to the Active Ingredients in Robitussin[®] DM**

Adult Tussin DM

Cough Suppressant/ Expectorant

Dextromethorphan HBr 20 mg per 20 mL

Guaifenesin 200 mg per 20 mL

For Ages 12 & over

Helps Control Cough

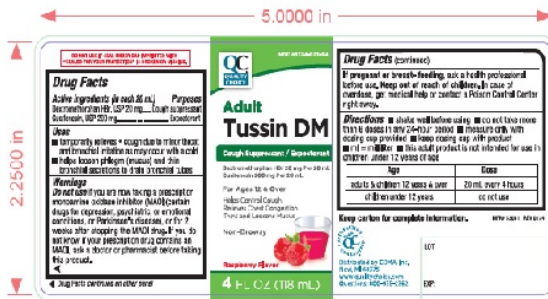
Relieves Chest Congestion

Thins and Loosens Mucus

Non- Drowsy

Raspberry Flavor

4 FL OZ (118 mL)





NDC 83324-023-08

QUALITY CHOICE

***Compare to the Active Ingredients in Robitussin ® DM**

Adult Tussin DM

Cough Suppressant/ Expectorant

Dextromethorphan HBr 20 mg per 20 mL

Guaifenesin 200 mg per 20 mL

For Ages 12 & over

Helps Control Cough

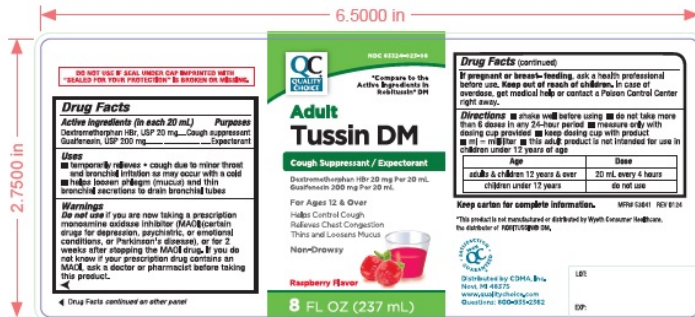
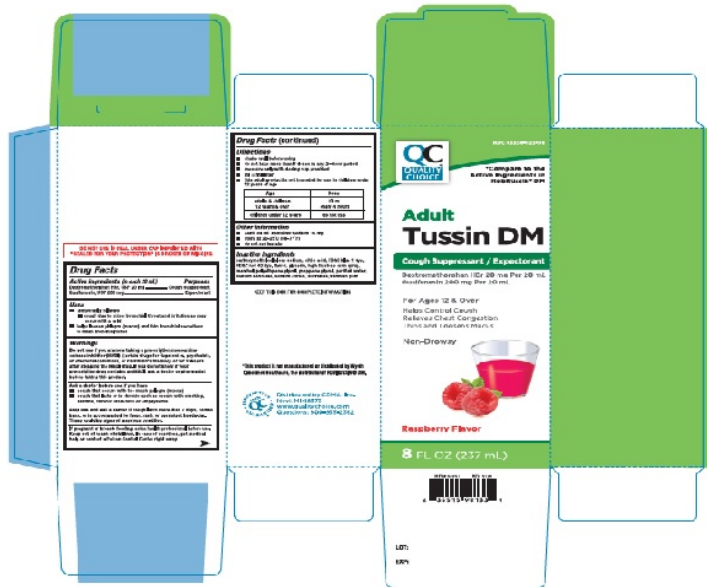
Relieves Chest Congestion

Thins and Loosens Mucus

Non- Drowsy

Raspberry Flavor

8 FL OZ(237 mL)



DO NOT USE IF SEAL UNDER CAP IMPRINTED WITH "SCALD FOR YOUR PROTECTION" IS BROKEN OR UNUSUAL.

Drug Facts

Active Ingredients (In each 20 mL) Purpose
 Dextromethorphan HBr, USP 20 mg...Cough suppressant
 Guaifenesin, USP 200 mg...Expectorant

Usage
 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 if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, 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.

Drug Facts continued on other panel

QC QUALITY CHOICE

Compare to the Active Ingredients in Robitussin® DM

Adult Tussin DM

Cough Suppressant / Expectorant

Dextromethorphan HBr 20 mg Per 20 mL
 Guaifenesin 200 mg Per 20 mL

For Ages 12 & Over
 Helps Control Cough
 Relieves Chest Congestion
 Thins and Loosens Mucus
 Non-Drowsy

Raspberry Flavor

8 FL OZ (237 mL)

Drug Facts (continued)

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 Shake well before using. Do not take more than 6 doses in any 24-hour period. Measure only with dosing cup provided. Keep dosing cup with product. 10 mL = 2 teaspoons. This adult product is not intended for use in children under 12 years of age.

Age	Dose
adults & children 12 years & over	20 mL every 4 hours
children under 12 years	do not use

Keep carton for complete information. NDC 43334-0234-8
 This product is manufactured or distributed by West Consumer Health, the exclusive of FORTICOR INC.

QC logo
 Distributed by CSM, Inc., New, NH 03275
 www.westconsumerhealth.com
 Q1001011-0000000262

LOT:
 EXP:

QCH ADULT TUSSIN DM 542

dextromethorphan hbr, guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-023
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 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
HIGH FRUCTOSE CORN SYRUP (UNII: XY6UN3QB6S)	
MENTHOL (UNII: L7T10EIP3A)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	RED	Score	
Shape		Size	
Flavor	RASPBERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-023-04	1 in 1 CARTON	05/08/2024	
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

2	NDC:83324-023-08	1 in 1 CARTON	05/08/2024	
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 Drug	part341	05/08/2024	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Establishment

Name	Address	ID/FEI	Business Operations
Guardian Drug Company		119210276	MANUFACTURE(83324-023)

Revised: 5/2024

Chain Drug Marketing Association Inc.