QCH ADULT TUSSIN DM SUGAR FREE 545 - dextromethorphan hbr, guaifenesin liquid
Chain Drug Marketing Association Inc.

QCH Adult Tussin DM Sugar Free 545

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 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 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

- 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 20-25ºC (68-77ºF)
- do not refrigerate

INACTIVE INGRADIENTS

carboxymethylcellulose sodium, citric acid, flavor, glycerin, menthol, polyethylene glycol, propylene glycol, purified water , sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum.

PRINCIPAL DISPLAY PANEL

NDC 83324-025-04

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

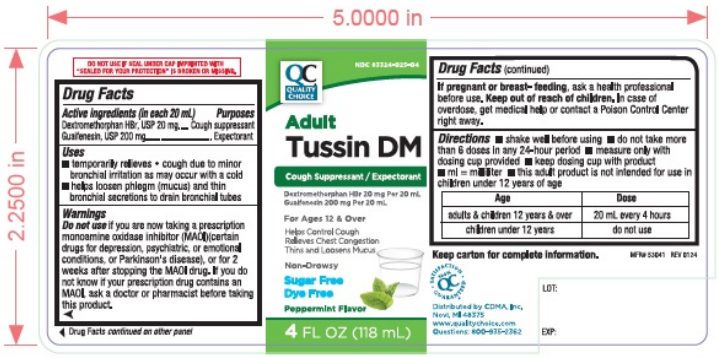
Specially Formulated for Diabetics

Non-Drowsy

Sugar Free Dye Free

Peppermint Flavor

4 FL OZ (118 mL)



QCH ADULT TUSSIN DM SUGAR FREE 545

dextromethorphan hbr, guaifenesin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-025
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)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL (UNII: L7T10EIP3A)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	PEPPERMINT	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-025-04	1 in 1 CARTON	05/08/2024	
1		118 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(83324-025)

Revised: 5/2024 Chain Drug Marketing Association Inc.