QCH MAXIMUM STRENGTH MUCUS RELIEF 617- dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid
Chain Drug Marketing Association Inc.

QCH Maximum Strength Mucus Relief 617

ACTIVE INGREDIENTS (in each 20 mL)

Dextromethorphan HBr 20 mg
Guaifenesin 400 mg
Phenylephrine HCl 10 mg

PURPOSE

Cough suppressant

Expectorant

Nasal decongestant

USE(S)

• helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes and make coughs more productive
• temporarily relieves:

• cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants

• the intensity of coughing
• the impulse to cough to help you get to sleep
• nasal congestion due to a cold

WARNINGS

.

DO NOT USE

• for children 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

• heart disease
• high blood pressure
• thyroid disease
• diabetes
• trouble urinating due to an enlarged prostate gland
• persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema
• cough that occurs with too much phlegm (mucus)

WHEN USING THIS PRODUCT

• **do not use more than directed**

STOP USE AND ASK DOCTOR IF

- nervousness, dizziness or sleeplessness occur
- symptoms do not get better within 7 days or occur with fever
- cough comes back, or occurs with rash or persistent headache. These could be signs of a serious condition.

PREGNANCY/BREASTFEEDING

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
• measure only with dosing cup provided
• do not use dosing cup with other products
• dose as follows or as directed by a doctor
• **Adults and children 12 years and olde**r: 20 ml in dosing cup provided every 4 hours .
• **Children under 12 years of age**: Do not use

OTHER INFORMATION

• **each 20 ml contains**: sodium 20 mg
• store between 15-30ºC (59-86ºF)
• do not refrigerate
• dosing cup provided

INACTIVE INGREDIENTS

citric acid anhydrous, edetate disodium, FD&C blue #1, FD&C red #40, flavors, glycerin, propylene glycol, propyl gallate, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum.

PRINCIPAL DISPLAY PANEL

NDC 83324-144-06

QC® QUALITY CHOICE

***Compare to the Active Ingredients in Maximum Strength Mucinex® Fax Max™ Severe Congestion & Cough**

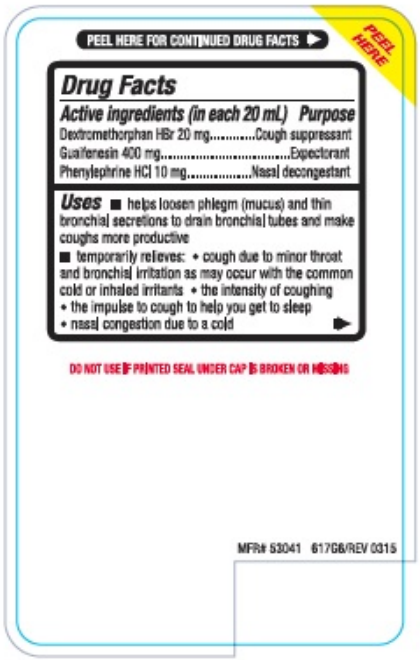
Maximum Strength Mucus Relief Sever Congestion & Cough

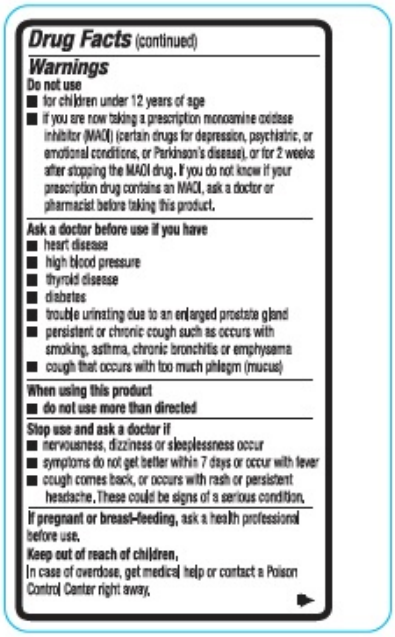
Dextromethorphan HBr 20 mg - Cough suppressant
Guaifenesin 400 mg - Expectorant
Phenylephrine HCl 10 mg - Nasal decongestant

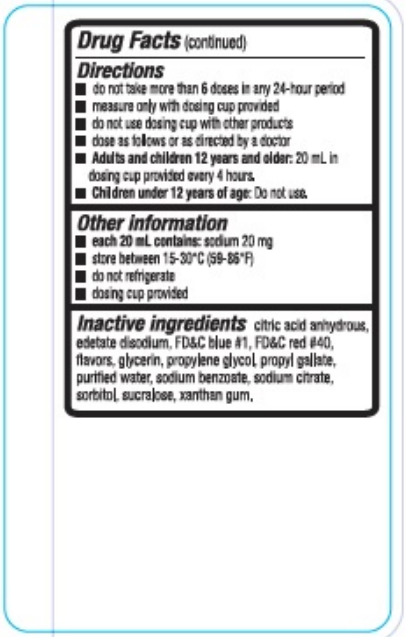
Helps Control Cough
Relieves Nasal & Chest Congestion
Thins & Loosens Mucus

6 FL OZ (177 mL)


NDC 83324-144-06
QC QUALITY CHOICE
*Compare to the Active Ingredients in Maximum Strength Mucinex® Fast Max™ Severe Congestion & Cough
Maximum Strength
Mucus Relief
Severe Congestion & Cough
Dextromethorphan HBr - Cough Suppressant
Guaifenesin - Expectorant
Phenylephrine HCl - Nasal Decongestant
Helps Control Cough
Relieves Nasal & Chest Congestion
Thins & Loosens Mucus
6 FL OZ (177 mL)
Distributed by CDMA, Inc.
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362
MFR# 53041 REV 0724
*This product is not manufactured or distributed by Reckitt Benckiser, the distributor of Mucinex® Fast Max™ Severe Congestion and Cough.
6 35515 97024 3


PEEL HERE FOR CONTINUED DRUG FACTS
PEEL HERE
Drug Facts
Active ingredients (in each 20 mL) Purpose
Dextromethorphan HBr 20 mg.....Cough suppressant
Guaifenesin 400 mg.....Expectorant
Phenylephrine HCl 10 mg.....Nasal decongestant
Uses ■ helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes and make coughs more productive
■ temporarily relieves: • cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants • the intensity of coughing • the impulse to cough to help you get to sleep • nasal congestion due to a cold
DO NOT USE IF PRINTED SEAL UNDER CAP IS BROKEN OR MISSING
MFR# 53041 617GB/REV 0315


Drug Facts (continued)
Warnings
Do not use
■ for children 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
■ heart disease
■ high blood pressure
■ thyroid disease
■ diabetes
■ trouble urinating due to an enlarged prostate gland
■ persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema
■ cough that occurs with too much phlegm (mucus)
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 occur with fever
■ cough comes back, or occurs with 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.


Drug Facts (continued)
Directions
■ do not take more than 6 doses in any 24-hour period
■ measure only with dosing cup provided
■ do not use dosing cup with other products
■ dose as follows or as directed by a doctor
■ Adults and children 12 years and older: 20 mL in dosing cup provided every 4 hours.
■ Children under 12 years of age: Do not use.
Other information
■ each 20 mL contains: sodium 20 mg
■ store between 15-30°C (59-86°F)
■ do not refrigerate
■ dosing cup provided
Inactive ingredients citric acid anhydrous, edetate disodium, FD&C blue #1, FD&C red #40, flavors, glycerin, propylene glycol, propyl gallate, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum.

QCH MAXIMUM STRENGTH MUCUS RELIEF 617

dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83324-144
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	400 mg in 20 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	BLUE	**Score**	
Shape		**Size**	
Flavor	BERRY (mixed berry)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-144-06	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/06/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/06/2024	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Establishment

Name	Address	ID/FEI	Business Operations
Guardian Drug Company		119210276	MANUFACTURE(83324-144)

Revised: 9/2024 Chain Drug Marketing Association Inc.