

MUCUS RELIEF DM HONEY AND BERRY FLAVOR- dextromethorphan hydrobromide and guaifenesin solution
RARITAN PHARMACEUTICALS INC

DRx Choice® Mucus Relief DM Honey & Berry Flavor

Drug Facts

<i>Active ingredients (in each 20 mL)</i>	<i>Purposes</i>
Dextromethorphan HBr 20 mg	Cough suppressant
Guaifenesin 400 mg	Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus 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

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

- 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

- cough lasts more than 7 days, comes back, or occurs with fever, rash, or headache that lasts. 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 at 1-800-222-1222.

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 of age and over:** 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 8 mg
- store at room temperature
- do not refrigerate

Inactive ingredients

citric acid, edetate disodium, flavors, glycerin, propylene glycol, purified water, sodium benzoate, sorbitol, sucralose, xanthan gum

Questions?

1-866-467-2748

PRINCIPAL DISPLAY PANEL - 180 mL Bottle Label

*Compare to the active ingredient in Maximum Strength Mucinex® Fast Max® DM Max Honey & Berry Flavor

NDC 68163-605-06

DRx Choice

maximum Strength

Mucus

relief DM

Dextromethorphan HBr – Cough Suppressant

Guaifenesin – Expectorant

- Controls Cough
- Relieves Chest Congestion
- Thins & Loosens Mucus
- 4 Hour Dosing

Honey& Berry Flavor

Naturally and Artificially Flavored

For Ages 12+

6 FL OZ (180mL)

Manufactured by:

Raritan Pharmaceuticals

8 Joanna Court

East Brunswick,

NJ 08816

*This product is not manufactured or distributed by RB Health (US) the distributor of Maximum Strength Mucinex® Fast-Max® DM Max Honey & Berry Flavor



MUCUS RELIEF DM HONEY AND BERRY FLAVOR

dextromethorphan hydrobromide and guaifenesin solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68163-605
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	HONEY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68163-605-06	180 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package	04/21/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	04/21/2023	

Labeler - RARITAN PHARMACEUTICALS INC (127602287)

Revised: 9/2025

RARITAN PHARMACEUTICALS INC