MAXIMUM STRENGTH MUCUS RELIEF DM- dextromethorphan hydrobromide and guaifenesin solution
CARDINAL HEALTH

LEADER Maximum Strength Mucus Relief DM 6 FL OZ

Drug Facts

Active ingredients (in each 20 mL)	***Purposes***
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

- 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

- 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 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 at 1-800-222-2222.

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
- mL = milliliter
- **Adults and children 12 years and older:** 20 mL 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
- dosing cup provided
- low sodium

Inactive ingredients

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

Questions or comments?

1-866-467-2748

PRINCIPAL DISPLAY PANEL - 180 mL Bottle Label

LEADER

NDC 70000-0129-1

Maximum Strength

Mucus Relief DM

Dextromethorphan HBr | Guaifenesin
Cough Suppressant | Expectorant

COMPARE TO MAXIMUM STRENGTH MUCINEX® FAST MAX® DM MAX active ingredients*

100% Money back Guarantee

For Ages 12+

- **Controls Cough**
- **Relieves Chest Congestion**
- **Thins & Loosens Mucus**
- **4 Hour Dosing**
- **Maximum Strength Formula**

6 FL OZ (180 mL)

Tamper evident: do not use if printed seal under cap is broken or missing.

DISTRIBUTED BY CARDINAL HEALTH

DUBLIN, OHIO 43017

www.myleader.com

1-800-200-6313

Essential to care™ since 1979

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Maximum Strength Mucinex® Fast -Max® DM Max.

MAXIMUM STRENGTH MUCUS RELIEF DM

dextromethorphan hydrobromide and guaifenesin solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70000-0129
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

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)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
propylene glycol (UNII: 6DC9Q167V3)	
propyl gallate (UNII: 8D4SNN7V92)	
water (UNII: 059QF0KO0R)	
sodium benzoate (UNII: OJ245FE5EU)	
sorbitol (UNII: 506T60A25R)	
sucralose (UNII: 96K6UQ3ZD4)	
xanthan gum (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0129-1	180 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/06/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/06/2017	

Labeler - CARDINAL HEALTH (063997360)

Revised: 11/2024 CARDINAL HEALTH