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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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 final	part341	03/06/2017	

Labeler - CARDINAL HEALTH (063997360)

Revised: 10/2023 CARDINAL HEALTH