

**MUCUS RELIEF DM MAXIMUM STRENGTH- dextromethorphan hbr,  
guaifenesin liquid  
P & L Development, LLC**

*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.*

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**Drug Facts**

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

Dextromethorphan HBr 20 mg

Guaifenesin 400 mg

**Purposes**

Cough suppressant

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 (1800-222-1222) right away.

## **Directions**

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device
- keep dosing cup with product
- mL = milliliter
- dose as follows or as directed by a doctor
- adults and children 12 years of age and older: 20 mL every 4 hours
- children under 12 years of age: do not use

## **Other information**

- each 20 mL contains: sodium 20 **mg**
- store between 20-25°C (68-77°F). Do not refrigerate.

## **Inactive ingredients**

anhydrous citric acid, disodium EDTA FD&C red #40, flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum

## **Principal Display Panel**

\*Compare to the active ingredients in Maximum Strength Mucinex® Fast-Max® DM Max maximum strength mucus Relief dm max

Dextromethorphan HBr

Guaifenesin

relieves

- cough
- Chest congestion & mucus
- 4-hour dosing

for ages 12 years & over

FL OZ (mL)

\*This product is not manufactured or distributed by Reckitt Benckiser, distributor of Maximum Strength Mucinex® Fast-Max® DM Max

**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND OR UNDER CAP IS BROKEN OR MISSING.**

**Manufactured by:**

**PL Developments**

**11865 S. Alameda St**

**Lynwood, CA 90262**

**Questions or comments? Call 1-877-753-3935 Monday-Friday 9AM-5PM EST**

**Package Label**



\*Compare to the active ingredients in  
Maximum Strength Mucinex®  
Fast-Max® DM Max  
NDC 49580-0505-6

maximum strength  
**mucus relief**  
**dm max**  
dextromethorphan HBr  
guaifenesin

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6 fl oz (177 mL)



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**PARENTS:**  
Learn about teen medicine abuse  
[www.StopMedicineAbuse.org](http://www.StopMedicineAbuse.org)

PLD-D409A  
LB004108

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Guaifenesin 400 mg	Expectorant

PEEL CORNER FOR MORE DRUG FACTS

**Drug Facts (continued)**

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PEEL CORNER FOR MORE DRUG FACTS

**ReadyinCase Mucus Relief DM Max**

# MUCUS RELIEF DM MAXIMUM STRENGTH

dextromethorphan hbr, guaifenesin liquid

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:49580-0505
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>PROPYL GALLATE</b> (UNII: 8D45NN7V92)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	

## Packaging

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:49580-0505-6	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/30/2016	

## Marketing Information

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC monograph final	part341	04/30/2016	

**Labeler** - P & L Development, LLC (101896231)

