

**MUCUS RELIEF DM- guaifenesin, dextromethorphan hbr tablet**  
**P & L Development, LLC**

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**Drug Facts**

**Active ingredients (in each extended-release tablet)**

Dextromethorphan HBr 30 mg

Guaifenesin 600 mg

**Purpose**

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**

- 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 accompanied by 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 illness.

**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 (1-800-222-1222).

**Directions**

- do not crush, chew, or break tablet
- take with a full glass of water
- this product can be administered without regards for timing of meals
- adults and children 12 years of age and older: 1 or 2 tablet every 12 hours; not more than 4 tablets in 24 hours
- children under 12 years of age: do not use

**Other information**

store between 20° to 25°C (68° to 77°F)

**Inactive ingredients**

carbomer, colloidal silicon dioxide, D&Cyellow #10 aluminum lake, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, povidone, talc

**Questions or comments?**

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

**Principal Display Panel**

Compare to active ingredients in Mucinex® DM\*

mucus relief dm

guaifenesin 600 mg

expectorant

dextromethorphan HBr

cough suppressant

- 12-hour relief
- Controls Cough
- Thins & Loosens Mucus

extended-release tablets

\*This product is not manufactured or distributed Reckitt Benckiser LLC, distributor of Mucinex® DM.

**TAMPER EVIDENT: DO NOT USE IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.**

**KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.**

Distributed by:  
PL Developments  
200 Hicks Street  
Westbury, NY 11590

### Package Label

<b>Drug Facts</b>	
<b>Active ingredients (in each extended-release tablet)</b> Dextromethorphan HBr 30 mg Guaifenesin 600 mg	<b>Purposes</b> Cough Suppressant Expectorant
<b>Uses</b> ■ 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	
<b>Warnings</b> <b>Do not use</b> ■ 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. <b>Ask a doctor before use if you have</b> ■ persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema ■ cough accompanied by too much phlegm (mucus) <b>When using this product,</b> do not use more than directed. <b>Stop use and ask a doctor if</b> cough lasts more than 7 days, comes back, or occurs with fever, rash or persistent headache. These could be signs of a serious illness. If pregnant or breast-feeding, ask a health professional before use. <b>Keep out of the reach of children.</b> In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).	
<b>Directions</b> ■ do not crush, chew, or break tablet ■ take with a full glass of water ■ this product can be administered without regard for timing of meals ■ adults and children 12 years of age and older: 1 or 2 tablets every 12 hours; not more than 4 tablets in 24 hours ■ children under 12 years of age: do not use	
<b>Other information</b> ■ store between 20° to 25°C (68° to 77°F)	
<b>Inactive ingredients</b> carbomer, colloidal silicon dioxide, D&C yellow #10 aluminum lake, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, povidone, talc	
<b>Questions or comments?</b> Call 1-877-753-3835 Monday-Friday 9AM-5PM EST	

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Distributed by:  
**PL Developments**  
200 Hicks Street  
Westbury, NY 11590



Compare to active ingredients in  
**Mucinex® DM\***  
NDC 59726-733-20

mucus relief dm

mucus relief dm

guaifenesin 600 mg  
expectorant  
dextromethorphan HBr 30 mg  
cough suppressant

- 12-hour relief
- controls cough
- thins and loosens mucus

20 extended-release tablets



actual size

TAMPER EVIDENT: DO NOT USE IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Lot No.:  
Exp. Date:

**READYinCASE Mucus Relief DM**

**MUCUS RELIEF DM**

guaifenesin, dextromethorphan hbr tablet

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:59726-733
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	600 mg
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg

## Inactive Ingredients

<b>Ingredient Name</b>	<b>Strength</b>
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>CARBOMER 934</b> (UNII: Z135WT9208)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>POVIDONE</b> (UNII: FZ989GH94E)	
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	

## Product Characteristics

<b>Color</b>	yellow	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	16mm
<b>Flavor</b>		<b>Imprint Code</b>	AN038
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59726-733-40	40 in 1 CARTON	12/31/2018	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:59726-733-06	6 in 1 CARTON	12/31/2018	
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:59726-733-20	20 in 1 CARTON	12/31/2018	
3		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA209692	12/31/2018	

**Labeler** - P & L Development, LLC (800014821)

Revised: 1/2026

P & L Development, LLC