

# **MUCINEX DM- guaifenesin and dextromethorphan hydrobromide tablet, extended release**

## **Select Consumer Group**

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**Mucinex®DM**

### **Drug Facts**

Active ingredients (in each extended-release tablet)

Dextromethorphan HBr 30 mg

Guaifenesin 600 mg

<b>Active ingredients (in each extended-release tablet)</b>	<b>Purposes</b>
Dextromethorphan HBr 30 mg	Cough suppressant
Guaifenesin 600 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 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.

### **Directions**

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

### **Other information**

- store at 20-25°C (68-77°F)

### **Inactive ingredients**

carbomer homopolymer type B; D&C yellow no. 10 aluminum lake; hypromellose, USP; magnesium stearate, NF; microcrystalline cellulose, NF; sodium starch glycolate, NF

### **Questions?**

**1-866-MUCINEX (1-866-682-4639)**

Dist. by: RB Health (US)  
Parsippany, NJ 07054-0224

Made in England

### **PRINCIPAL DISPLAY PANEL - 20 Tablet Blister Pack Carton**

NDC 63824-056-32

Mucinex®DM

600 mg guaifenesin & 30 mg dextromethorphan HBr  
extended-release tablets

EXPECTORANT & COUGH SUPPRESSANT

12  
HOUR®

- Controls Cough
- Thins and Loosens Mucus
- Immediate and Extended Release

20  
EXTENDED-RELEASE TABLETS



## MUCINEX DM

guaifenesin and dextromethorphan hydrobromide tablet, extended release

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:85237-1650(NDC:63824-056)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	600 mg
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED)</b> (UNII: HHT01ZNK31)	
<b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)	
<b>ALUMINUM OXIDE</b> (UNII: LMI26O6933)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	

<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	

### Product Characteristics

<b>Color</b>	white (yellow and white)	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	16mm
<b>Flavor</b>		<b>Imprint Code</b>	Mucinex;600
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85237-1650-1	2 in 1 POUCH; Type 0: Not a Combination Product	12/27/2025	
2	NDC:85237-1650-2	1 in 1 BLISTER PACK	12/24/2025	
2		2 in 1 POUCH; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA021620	06/26/2012	

**Labeler** - Select Consumer Group (119185084)

**Registrant** - Select Consumer Group (119185084)

### Establishment

Name	Address	ID/FEI	Business Operations
Select Consumer Group		119185084	repack(85237-1650)

Revised: 12/2025

Select Consumer Group