

**MUCUS RELIEF DM EXTENDED RELEASE CAPLETS- guaifenesin,  
dextromethorphan hbr tablet, extended release  
ATLANTIC BIOLOGICALS CORP.**

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**Mucus Relief DM Extended Release**

**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

**Description**

Mucus Relief DM ER

Guaifenesin 600 mg (Expectorant)

Dextromethorphan HBr 30 mg (Cough Suppressant)

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

Extended-release tablets, compared to the active ingredients in Mucinex DM.

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

**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 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 (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?**

**DISTRIBUTED BY:**

**ATLANTIC BIOLOGICALS CORP.**

20101 NE 16TH PL

MIAMI, FL 33179

Call 1-800-509-7592

Monday-Friday 9 AM - 5 PM EST

### Package Label

17856-1162-01

Mucinex DM

(Guaifenesin/

Dextromethorphan HBr)

DELIVERS

30-600mg/1 ER TAB



See package insert for indications and dosage schedule

Helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and makes coughs more productive. DO NOT Break, Chew, or Crush tablet. Store at 20°-25°C (68°-77°F)  
\*\*Keep this and all Medications out of the reach of children\*\*



17856-1162-01 Dosage 30-600mg/1 TAB

Mucus Relief (Guaifenesin/  
Dextromethorphan HBr)

Qty: 100 ER TABLETS



GTIN: 00317856116213

S/N: XXXXXXXXXXXX

Exp: 03/25/25

Lot: XXXXXXXXXXXX



Distributed by: Atlantic Biologicals Corp.  
Miami, FL 33179

Rev.08/21

Call to Reorder:

## MUCUS RELIEF DM EXTENDED RELEASE CAPLETS

guaifenesin, dextromethorphan hbr tablet, extended release

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:17856-1162(NDC:0536-1161)
<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: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg

## Inactive Ingredients

Ingredient Name	Strength
<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:17856-1162-1	100 in 1 CARTON	03/17/2025	
1		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	12/13/2018	

**Labeler** - ATLANTIC BIOLOGICALS CORP. (047437707)

**Registrant** - ATLANTIC BIOLOGICALS CORP. (047437707)