

**MUCUS RELIEF DM IMMEDIATE RELEASE- mucus relief dm tablet**  
**Allegiant Health**

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**432 - Mucus Relief DM**

**Active ingredient(s)**

Dextromethorphan HBr 20mg  
Guaifenesin 400mg

**Purpose**

Cough suppressant  
Expectorant

**Use(s)**

- 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**

- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema
- cough accompanied by too much phlegm (mucus)

**Ask a doctor or pharmacist before use if**

taking sedatives or tranquilizers

### **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

### **Pregnancy/Breastfeeding**

If pregnant or breast-feeding, ask a health professional before use.

### **Keep out of reach of children**

In case of accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

### **Directions**

**Adults and children 12 years and older:** take 1 tablet every 4 hours with a full glass of water while symptoms persist. Do not exceed 6 doses in 24 hours.

**Children under 12 years of age:** do not use

### **Other information**

- store at 25°C (77°F) excursions between 15°-30°C (59°-86°F)
- keep in a dry place and do not expose to heat
- do not use if imprinted safety seal under cap is broken or missing
- You may report side effects to 1-888-952-0050

### **Inactive ingredients**

croscarmellose sodium, magnesium stearate, maltodextrin, microcrystalline cellulose, povidone, silicon dioxide, stearic acid

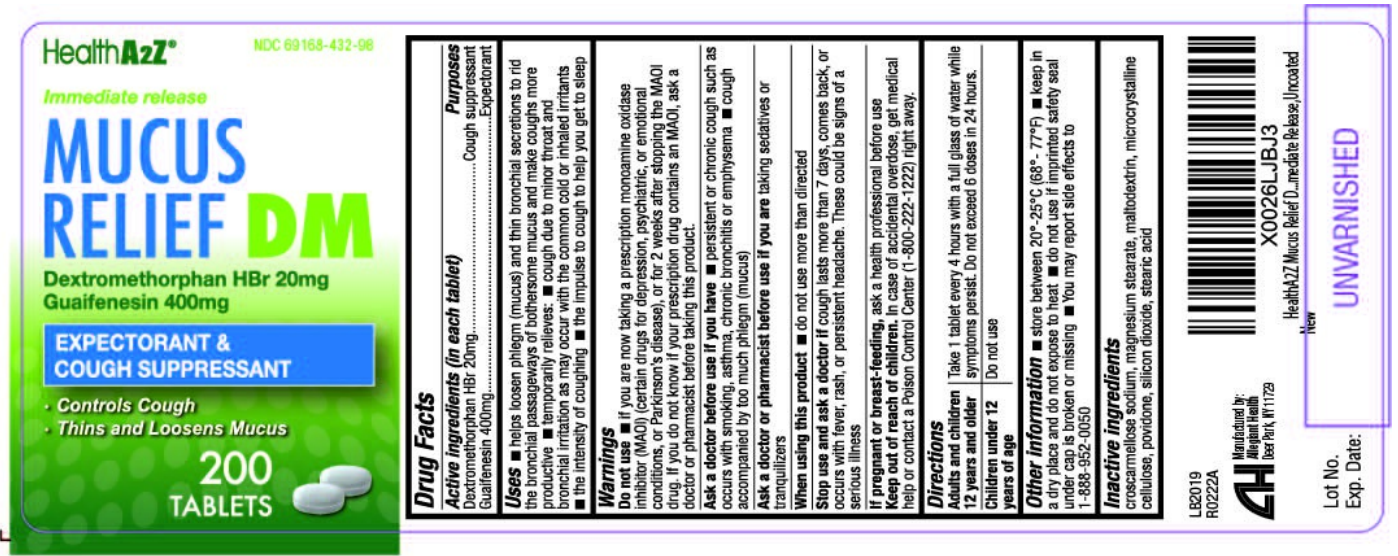
### **Questions**

Questions or comments?

Call 1-888-952-0050

Monday through Friday 9AM- 5PM EST.

### **Principal Display Panel**



## Mucus Relief DM

### MUCUS RELIEF DM IMMEDIATE RELEASE

mucus relief dm tablet

#### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69168-432
<b>Route of Administration</b>	ORAL		

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg

#### Inactive Ingredients

Ingredient Name	Strength
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POVIDONE</b> (UNII: FZ989GH94E)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	

#### Product Characteristics

<b>Color</b>	white	<b>Score</b>	2 pieces
<b>Shape</b>	OVAL	<b>Size</b>	16mm

<b>Flavor</b>		<b>Imprint Code</b>	AH432;bisect	
<b>Contains</b>				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69168-432-98	200 in 1 BOTTLE; Type 0: Not a Combination Product	05/18/2022	
2	NDC:69168-432-07	1 in 1 CARTON	05/18/2022	
2		10 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:69168-432-26	1 in 1 CARTON	05/18/2022	
3		20 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:69168-432-50	50 in 1 BOTTLE; Type 0: Not a Combination Product	06/08/2022	
5	NDC:69168-432-06	120 in 1 BOTTLE; Type 0: Not a Combination Product	11/06/2023	
6	NDC:69168-432-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	11/06/2023	
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M012	05/18/2022		

**Labeler** - Allegiant Health (079501930)

Revised: 5/2022

Allegiant Health