

**MUCINEX MAXIMUM STRENGTH- guaifenesin tablet, extended release**  
**A-S Medication Solutions**

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**MUCINEX MAXIMUM STRENGTH**

***Drug Facts***

**Active ingredient (in each extended-release bi-layer tablet)**

Guaifenesin 1200 mg

**Purpose**

Expectorant

**Uses**

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

**Warnings**

**Do not use**

- for children under 12 years of age

**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)

**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 tablet
- take with a full glass of water
- this product can be administered without regard for the timing of meals
- adults and children 12 years of age and over: 1 tablet every 12 hours. Do not exceed 2 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; FD&C blue no. 1 aluminum lake; hypromellose, USP; magnesium stearate, NF; microcrystalline cellulose, NF; sodium starch glycolate, NF

## Questions?

**1-866-MUCINEX (1-866-682-4639)** You may also report side effects to this phone number.

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

Made in England

## HOW SUPPLIED

Product: 50090-2301

NDC: 50090-2301-0 7 TABLET, EXTENDED RELEASE in a BLISTER PACK / 1 in a CARTON

## Guaifenesin



## MUCINEX MAXIMUM STRENGTH

guaifenesin tablet, extended release

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50090-2301(NDC:63824-023)
<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	1200 mg

**Inactive Ingredients**

Ingredient Name	Strength
<b>Carbomer Homopolymer Type B (Allyl Pentaerythritol Crosslinked)</b> (UNII: HHT01ZNK31)	
<b>FD&amp;C blue no. 1</b> (UNII: H3R47K3TBD)	
<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 (blue and white)	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	22mm
<b>Flavor</b>		<b>Imprint Code</b>	Mucinex;1200
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090-2301-0	1 in 1 CARTON	02/11/2016	
1		7 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
NDA	NDA021282	06/26/2012	

**Labeler** - A-S Medication Solutions (830016429)**Establishment**

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-2301)