

MUCUS EXTENDED RELEASE- guaifenesin tablet, extended release
AMERISOURCEBERGEN DRUG CORPORATION

Mucus
Extended Release

Drug Facts

Active ingredient (in each extended-release 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 (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 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

- **Tamper evident: Do not use if carton is open or if printed seal on blister is broken or missing.**
- store between 20-25°C (68-77°F)

Inactive ingredients

carbomer homopolymer type B; hypromellose, USP; magnesium stearate, NF; microcrystalline cellulose, NF; sodium starch glycolate, NF

Questions?

(1-800-406-7984)

You may also report side effects to this phone number.

Distributed By:
AmerisourceBergen,
1300 Morris Drive
Chesterbrook, PA 19087

PRINCIPAL DISPLAY PANEL - 1200 mg Tablet Blister Pack Carton

NDC 46122-417-74

**Compare to active ingredient in
Maximum Strength Mucinex[®] †**

**GOOD
NEIGHBOR
PHARMACY[®]**

12 HOUR

Maximum Strength

Mucus • ER

**Guaifenesin Extended-Release
Tablets, 1200 mg**

Expectorant

- Relieves Chest Congestion
- Thins and Loosens Mucus
- Immediate and Extended Release

14 Extended-Release Tablets

Maximum Strength

Mucus • ER

Expectorant

Guaifenesin Extended-Release Tablets, 1200 mg

14 Extended-Release Tablets

NDC 46122-417-74

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Mucus • ER
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14 Extended-Release Tablets
Expectorant

Expectorant

14 Extended-Release Tablets

Maximum Strength
Mucus • ER
Guaifenesin Extended-Release Tablets, 1200 mg

Lot No.

Expiration Date:

NON VARNISH



5142025

Drug Facts

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Guaifenesin 1200 mg.....	Expectorant

Uses

Drug Facts (continued)

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Keep the carton. It contains important information. See end panel for expiration date.

[†]All trademarks are property of their respective owners. MUCINEX is a registered trademark of Rockitt Benckiser LLC.

GLUE - NO COATING

 Distributed By: AmerisourceBergen, 1300 Morris Drive, Chesterbrook, PA 19087. Visit us at www.gpcu.com. Made in England

ABC# 10177642

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0 87701 42951 4

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MUCUS EXTENDED RELEASE

guaifenesin tablet, extended release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46 122-417
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Guaifenesin (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ)	Guaifenesin	1200 mg

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: HHT01ZNK31)	
hypromellose, unspecified (UNII: 3NXW29V3WO)	
magnesium stearate (UNII: 70097M6I30)	

microcrystalline cellulose (UNII: OP1R32D61U)
sodium starch glycolate type A potato (UNII: 5856J3G2A2)

Product Characteristics

Color	WHITE	Score	no score
Shape	OVAL	Size	22mm
Flavor		Imprint Code	Mxeunic;1200
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46 122-417-74	1 in 1 CARTON	08/21/2017	
1		14 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	08/21/2017	

Labeler - AMERISOURCEBERGEN DRUG CORPORATION (007914906)

Establishment

Name	Address	ID/FEI	Business Operations
RECKITT BENCKISER HEALTHCARE INTERNATIONAL LTD		230780363	MANUFACTURE(46 122-417) , LABEL(46 122-417)

Revised: 11/2018

AMERISOURCEBERGEN DRUG CORPORATION