MUCUS RELIEF- guaifenesin 400 mg tablet Pioneer Life Sciences, LLC

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Active ingredient (in each tablet)

Guaifenesin 400 mg

Purpose

Expectorant

Uses

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

Warnings

Ask a doctor before use if you have

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

Stop use and ask a doctor if

• cough lasts for 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 care professional before use.

KEEP OUT OF REACH OF CHILDREN. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) immediately.

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: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
- Read all product information before using

Inactive ingredients

Colloidal Silicon Dioxide, Magnesium Stearate, Microcrystalline Cellulose, Stearic Acid Powder, Sodium Starch Glycolate

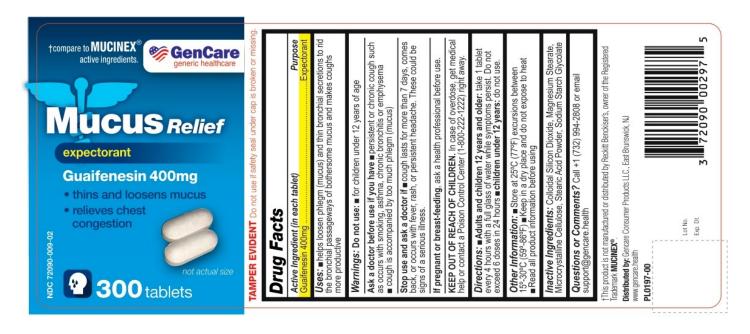
Questions or Comments?

Call +1-732-994-2808 or email support@gencare.health

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Distributed by: GenCare Consumer Products, LLC

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MUCUS RELIEF guaifenesin 400 mg tablet Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:72090-009 Route of Administration ORAL Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg

Inactive Ingredients	
Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics				
Color	white	Score	no score	
Shape	capsule	Size	17mm	
Flavor		Imprint Code	EB	
Contains				

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:72090-009- 02	300 in 1 BOTTLE; Type 0: Not a Combination Product	10/03/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/03/2024	

Labeler - Pioneer Life Sciences, LLC (014092742)

Revised: 10/2024 Pioneer Life Sciences, LLC