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

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**MUCUS RELIEF- Guaifenesin 400 mg tablet** 

#### 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

Do not use: for children under 12 years of age

#### Ask a doctor before use if you have

- persistent 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, Croscarmellose Sodium, Magnesium Stearate, Maize Starch, Microcrystalline Cellulose, Povidone K-30, Sodium Lauryl Sulphate

#### **Questions or Comments?**

Call 1-732-698-5070 Monday through Friday 9 am to 5 pm EST

This product is not manufcatured or distributed by Reckitt Benckiser's , owner of the Registered Trademark MUCINEX  $^{\circledR}$ 

**Distributed by:** GenCare Consumer Products, LLC 40E Cotters Ln Suite A, East Brunswick, NJ 08816

NDC 72090-013-01



# MUCUS RELIEF guaifenesin 400 mg tablet Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:72090-013 Route of Administration ORAL

## Active Ingredient/Active Moiety Ingredient Name Basis of Strength GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) GUAIFENESIN 400 mg

Inactive Ingredients	
Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
STARCH, CORN (UNII: O8232NY3SJ)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72090-013- 01	200 in 1 BOTTLE; Type 0: Not a Combination Product	01/19/2024	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M012	01/19/2024		

#### **Labeler - Pioneer Life Sciences, LLC (014092742)**

Revised: 1/2024 Pioneer Life Sciences, LLC