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

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

Ask a doctor before use if you have

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

When using this product

- do not exceed recommended dosage

Stop use and ask a doctor if

cough lasts for more than 7 days, recurs, or if accompanied by 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.

- **Do not crush, chew, or break tablet**
- **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 1-732-698-5070 Monday through Friday 9 am to 5 pm EST or www.pioneerlifesciences.com

This product is not manufcatured or distributed by Reckitt Benckiser's , owner of the Registered Trademark MUCINEX®

Distributed by: **GenCare Consumer Products, LLC**

40E Cotters Ln Suite A, East Brunswick, NJ 08816

NDC 72090-009-01

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:72090-009-01	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)

Registrant - Pioneer Life Sciences, LLC (014092742)

Revised: 1/2024 Pioneer Life Sciences, LLC