

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

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

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72090-044
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	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
POVIDONE K30 (UNII: U725QWY32X)				
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
Product Characteristics				
Color	white		Score	no score
Shape	capsule (Caplet)		Size	14mm
Flavor			Imprint Code	S400
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72090-044-01	200 in 1 BOTTLE; Type 0: Not a Combination Product	12/23/2025	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		M012	12/23/2025	

Labeler - Pioneer Life Sciences, LLC (014092742)

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