

MUCUS RELIEF IMMEDIATE RELEASE- guaifenesin 400mg tablet
TopCo Associates, LLC

370- Mucus Relief

Active ingredient(s)

Guaifenesin 400 mg

Purpose

Expectorant

Use(s)

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

Warnings

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 excessive phlegm (mucus)

When using this product

- do not exceed recommended dosage
- do not use for more than 7 days

Stop use and ask a doctor if

- cough lasts for more than 7 days, recurs, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.

Pregnancy/Breastfeeding

ask a healthcare professional before use.

Keep out of reach of children

In case of accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

Adults and children 12 years of age 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 of age: do not use

Other information

- do not use if imprinted safety seal under cap is broken or missing

Storage

- store at 25°C (77°F) excursions between 15° to 30°C (59° to 86°F)
- keep in a dry place and do not expose to heat

Inactive ingredients

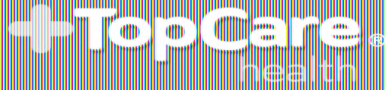
Croscarmellose Sodium, Maltodextrin, Microcrystalline Cellulose, Povidone, Silicon Dioxide, Stearic Acid

Questions

Call 1-888-423-0139

Principal Display Panel

NDC 76162-004-01



IMMEDIATE RELEASE

Mucus Relief

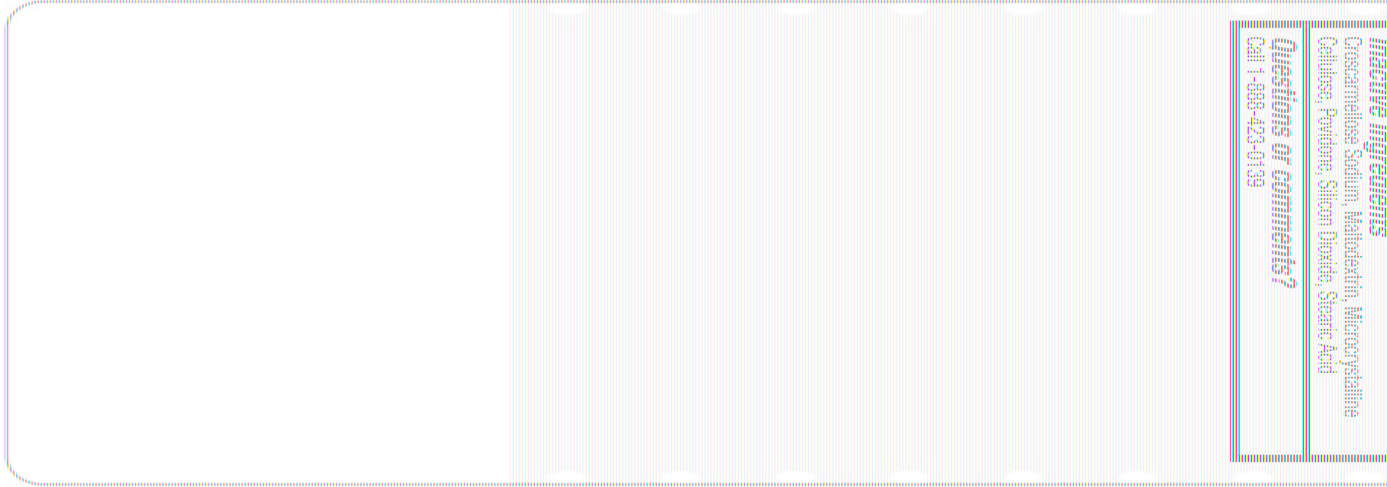
EXPECTORANT - GUAIFENESIN 400 mg

- Relieves Chest Congestion
- Thins & Loosens Mucus

100 TABLETS

Drug Facts	
Active ingredient (in each tablet) Guaifenesin 400 mg	Purpose Expectorant
Uses ■ helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive to rid the bronchial passageway of bothersome mucus	
Warnings 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 excessive phlegm (mucus) When using this product ■ do not exceed recommended dosage ■ do not use for more than 7 days Stop use and ask a doctor if ■ cough lasts for more than 7 days, recurs, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition.	
If pregnant or breast-feeding, ask a healthcare professional before use Keep out of reach of children. In case of accidental overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.	
<i>Drug Facts (continued on inside)</i>	

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✓ QUALITY GUARANTEED



Inactive ingredients Croscarmellose Sodium, Methocel K100, Microcrystalline Cellulose, Povidone, Silicon Dioxide, Stearic Acid
Questions or comments? Call 1-888-423-0139

Mucus Relief

MUCUS RELIEF IMMEDIATE RELEASE			
guaifenesin 400mg tablet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76162-004
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

CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
POVIDONE (UNII: FZ989GH94E)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	

Product Characteristics

Color	white	Score	2 pieces
Shape	OVAL	Size	17mm
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76162-004-81	2 in 1 CARTON	12/23/2014	01/31/2022
1		200 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:76162-004-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/23/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	12/23/2014	

Labeler - TopCo Associates, LLC (006935977)

Revised: 1/2021

TopCo Associates, LLC