MUCUS RELIEF- guaifenesin tablet Spirit Pharmaceuticals LLC

Mucus Relief

Drug Facts

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 make coughs more productive

Warnings

Do not use for children under 12 years of age

Ask a doctor before use if you have

• with persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema • cough accompanied by

Stop use and ask a doctor

if cough lasts more than 7 days, comes back, or occurs with fever, rash, or persistent headache. These could be signs of a serious illness.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

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

Directions

• do not crush, chew, or break tablet • take with a full glass of water • adults and children 12 years of age and over: take 1 tablet every 4 hours with a full glass of water while symptoms persist. Do not exceed 6 tablets in 24 hours. • children under 12 years of age: do not use

Other information

• store between 20-25°C (68-77°F)

Inactive ingredients

Colloidal silicon dioxide, magnesium stearate, maltodextrin, microcrystalline cellulose, polyvinyl pyrrolidone, sodium starch glycolate, stearic acid

Questions or comments?

1-888-333-9792

Distributed by:

Cabinet Health P.B.C.

Pouch





MUCUS RELIEF

guaifenesin tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68210-4173
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)			
MALTODEXTRIN (UNII: 7CVR7L4A2D)			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
PYRROLIDINE (UNII: LJU5627FYV)			
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)			
STEARIC ACID (UNII: 4ELV7Z65AP)			

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

F	Packaging			
#	tem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210- 4173-3	30 in 1 POUCH; Type 0: Not a Combination Product	11/19/2021	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	11/19/2021	

Labeler - Spirit Pharmaceuticals LLC (179621011)