MUCUS RELIEF- guaifenesin tablet SPIRIT PHARMACEUTICALS LLC

MUCUS RELIEF Caplets

Drug Facts

Active ingredient (in each caplet)

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

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

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 caplet
- take with a full glass of water
- adults and children 12 years of age and over: take 1 caplet every 4 hours with a full glass of water while symptoms persist. Do not exceed 6 caplets 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 sillicone dioxide, magnesium stearate, maltodextrin, microcrystalline cellulose, polyvinyl pyrrolidone, sodium starch glycolate, stearic acid

Questions or comments?

1-888-333-9792

PRINCIPAL DISPLAY PANEL

SOUNDBODY TM

MUCUS

RELIEF

Guaifenesin 400 mg

Expectorant

Thins &

Loosens

Mucus,

Relieves Chest Congestion

60 Caplets



MUCUS RELIEF

guaifenesin tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68210-0611

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)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

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

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210- 0611-6	1 in 1 CARTON	02/11/2020	
1		60 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	02/11/2020	

Labeler - SPIRIT PHARMACEUTICALS LLC (179621011)

Establishment

Name	Address	ID/FEI	Business Operations
ELYSIUM PHARMACEUTICALS LTD		915664486	manufacture(68210-0611)

Revised: 12/2023 SPIRIT PHARMACEUTICALS LLC