

**MUCUS RELIEF- guaifenesin tablet**  
**Prodose, Inc.**

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**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 condition.

**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 exceed 6 caplets in 24 hours.
- 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.
- 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?***

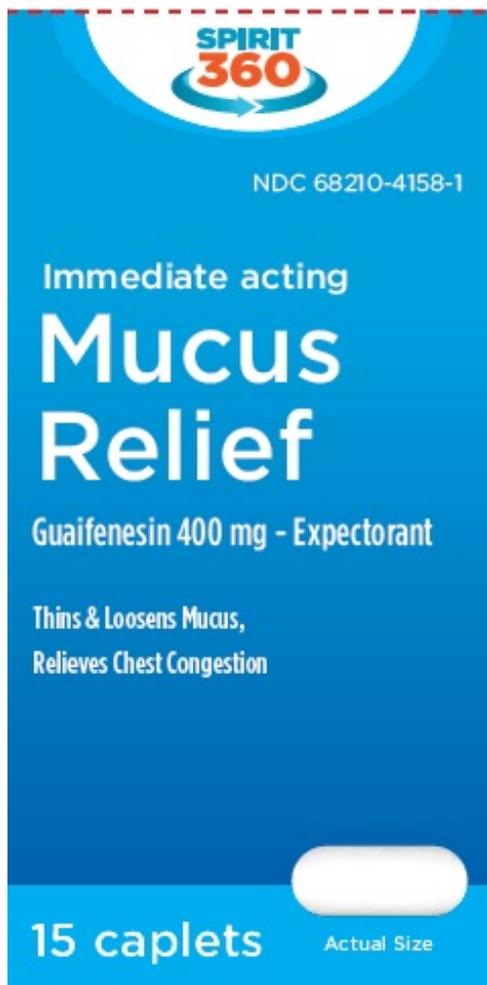
**1-888-333-9792**

### **PRINCIPAL DISPLAY PANEL**

Mucus Relief

Guaifenesin 400mg

- Expectorant
- Thins and Loosens Mucus
- Relieves Chest Congestion



## MUCUS RELIEF

guaifenesin tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68210-4158
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POVIDONE K30</b> (UNII: U725QWY32X)	
<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)	

STEARIC ACID (UNII: 4ELV7Z65AP)

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	17mm
<b>Flavor</b>		<b>Imprint Code</b>	EB
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210-4158-1	1 in 1 BOTTLE	05/25/2021	
1		15 in 1 BOTTLE; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/10/2020	

**Labeler** - Prodose, Inc. (119371190)

Revised: 1/2026

Prodose, Inc.