#### MUCUS RELIEF- guaifenesin tablet HealthLife of USA LLC

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#### Active ingredient (in each caplet)

Guaifenesin 400 mg

#### Purpose

Expectorant

#### Uses

• Temporary symptomatic relief from congested chests and cough.

#### Warnings

A persistent cough may be a sign of a serious condition. If cough persists for more than 1 week, tends to recur, or is accompanied by a fever, rash, or persistent headache, consult a doctor.

#### 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 consume alcohol, benzodiazepines or opioids.

## Stop use and ask a doctor if

cough lasts for more than 7 days please consult your physician or healthcare provider.

# 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 (1-800-222-1222) right away.

#### **Directions**

- Adults and children 12 years of age and over: take 1 caplet a day.
- Children under 12 years:Do not use.
- Do not exceed 4 doses in 24 hours.

#### Other information

- Store at 25°C (77°F) excursions between 15°-30°C (59°-86°F).
- Keep in a dry place and do not expose to heat.
- Read all product information before using.
- TEMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

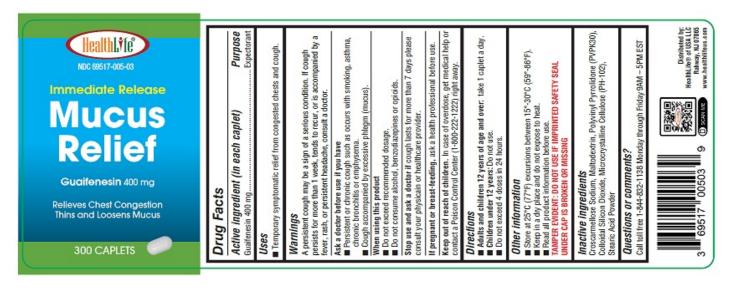
#### Inactive ingredients

Croscarmellose Sodium, Maltodextrin, Polyvinyl Pyrroliidone (PVPK30), Collidal Silicon Dioxide, Microcrystalline Cellulose (PH-102), Stearic Acid Powder

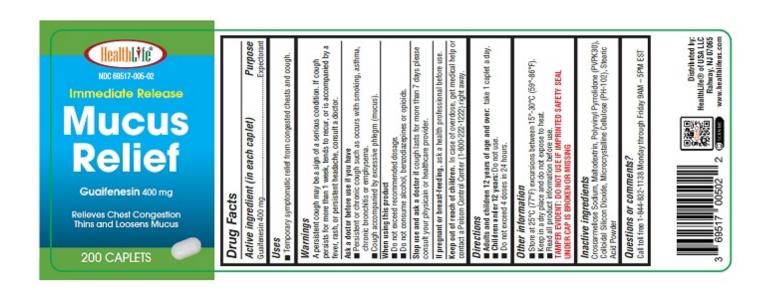
#### Questions or comments?

Call toll free 1-844-832-1138 Monday through Friday 9AM – 5PM EST

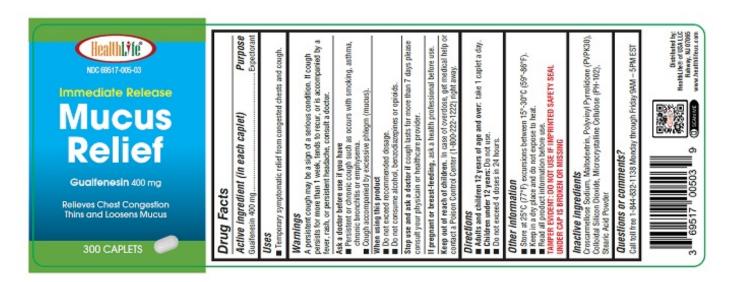
#### Principal Display Panel-60's Count



## Principal Display Panel-200's Count



### Principal Display Panel-300's Count



## **MUCUS RELIEF**

quaifenesin tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69517-005	
Route of Administration	ORAL			

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

## **Inactive Ingredients**

Ingredient Name	Strength
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POVIDONE K30 (UNII: U725QWY32X)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics				
Color	white	Score	no score	
Shape	CAPSULE	Size	14mm	
Flavor		Imprint Code	G400	
Contains				

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69517-005- 60	60 in 1 BOTTLE; Type 0: Not a Combination Product	09/26/2025			
2	NDC:69517-005- 02	200 in 1 BOTTLE; Type 0: Not a Combination Product	09/26/2025			
3	NDC:69517-005- 03	300 in 1 BOTTLE; Type 0: Not a Combination Product	09/26/2025			

Marketing Information				
Marketing Category				
OTC Monograph Drug	M012	09/26/2025		

# Labeler - HealthLife of USA LLC (079656178)

# Registrant - HealthLife of USA LLC (079656178)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
ELYSIUM PHARMACEUTICALS LIMITED		863182240	manufacture(69517-005)

Establishment					
Name	Address	ID/FEI	<b>Business Operations</b>		
HHH PHARMA USA LLC		062788820	pack(69517-005)		

Revised: 9/2025 HealthLife of USA LLC