# MUCUS RELIEF- guaifenesin tablet Safeway, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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#### **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 makes coughs more productive.

## Warnings

## Ask a doctor before use if you have

- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis or emphysema
- cough accopanied 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 (1-800-222-1222) right away.

#### **Directions**

- do not exceed 6 tablets in 24 hours
- take with a full glass of water

- adults and children 12 years of age and older
  - take 1 tablet every 4 hours 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, povidone, sodium starch glycolate, stearic acid

#### **Questions or comments?**

Call **1-888-723-3929** Monday-Friday 7AM-6PM PST

#### **Principal Display Panel**

#### **Mucus Relief**

GUAIFENESIN 400 mg TABLETS

**Expectorant** 

- Relieves Chest Congestion
- Thin & Loosens Mucus

#### **TABLETS**

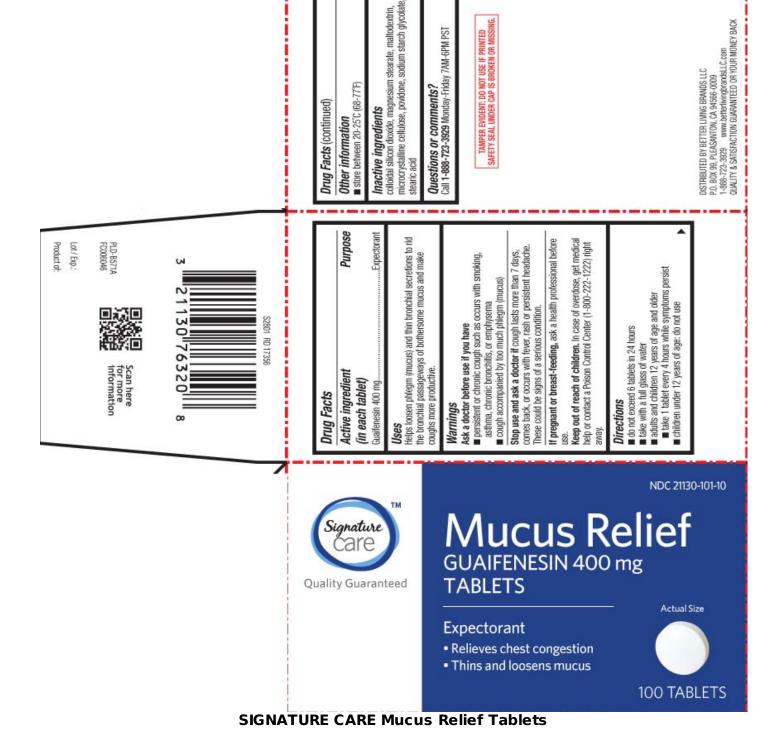
## TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

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P.O. BOX 99, PLEASANTON, CA 94566-0009

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## Package Label



## **MUCUS RELIEF**

guaifenesin tablet

	Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-101	
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			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
MALTODEXTRIN (UNII: 7CVR7L4A2D)				
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)				
POVIDONE (UNII: FZ989GH94E)				
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)				
STEARIC ACID (UNII: 4ELV7Z65AP)				

Product Characteristics				
Color	white	Score	2 pieces	
Shape	ROUND	Size	13mm	
Flavor		Imprint Code	TCL272	
Contains				

F	Packaging					
#	tem Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:21130- 101-10	1 in 1 BOX	08/31/2019	02/28/2025		
1		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product				

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part341	08/31/2019	02/28/2025	

## **Labeler -** Safeway, Inc. (009137209)

Revised: 11/2022 Safeway, Inc.