

**MUCUS RELIEF 600 MG - guaifenesin tablet, extended release**  
**MUCUS RELIEF MAXIMUM STRENGTH 1200 MG - guaifenesin tablet, extended release**  
**Guardian Drug Company**

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**Mucus Relief Guaifenesin Extended-Release Tablets 600 mg and 1200 mg**

**ACTIVE INGREDIENT(in each extended-release tablet)**

Guaifenesin 1200 mg

**PURPOSE**

Expectorant

**USE(S)**

helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

**WARNING**

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**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 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
- this product can be administered without regard for the timing of meals
- adults and children 12 years of age and over: 1 tablet every 12 hours.
- Do not exceed 2 tablets in 24 hours.
- children under 12 years of age: do not use

## **OTHER INFORMATION**

- tamper evident: do not use if carton is open or if printed seal on blister is broken or missing
- store between 20 to 25°C (68 to 77°F)

## **INACTIVE INGREDIENTS**

carbomer homopolymer, hypromellose, microcrystalline cellulose, povidone

## **QUESTIONS**

1-609-860-2600

Hours: 8am - 4pm, EST

You may also report side effects to this phone number.

## **PRINCIPAL DISPLAY PANEL**

NDC 53041-191-58

Compare to the active ingredient in Mucinex® Maximum Strength Extended Release  
1200 mg Tablets

GUARDIAN

12 HOUR

MAXIMUM STRENGTH

Mucus Relief

Guaifenesin Extended-Release Tablets 1200 mg

Expectorant

Relieves Chest Congestion

Thins and Loosens Mucus

14 EXTENDED-RELEASE TABLETS





## ACTIVE INGREDIENT (in each extended- release tablet)

Guaifenesin 600 mg

## PURPOSE

Expectorant

## USE(S)

helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

## WARNING

**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 tablet
- take with a full glass of water
- this product can be administered without regard for the timing of meals
- adults and children 12 years of age and over: 1 or 2 tablets every 12 hours.
- Do not exceed 4 tablets in 24 hours.
- children under 12 years of age: do not use

**OTHER INFORMATION**

- tamper evident: do not use if carton is open or if printed seal on blister is broken or missing
- store between 20 to 25°C (68 to 77°F)

**INACTIVE INGREDIENTS**

carbomer homopolymer, hypromellose, microcrystalline cellulose, povidone

**QUESTIONS?**

1-609-860-2600

Hours: 8am - 4pm, EST

You may also report side effects to this phone number.



## 20 EXTENDED-RELEASE TABLETS



## 500 EXTENDED-RELEASE TABLETS

NDC 53041-190-20

Compare to the active ingredient in Mucinex®  
Extended Release 600 mg Tablets\*

# Mucus Relief

*Guaifenesin Extended-Release  
Tablets 600 mg*

## Expectorant

Relieves Chest Congestion  
Thins and Loosens Mucus

actual size

**500** EXTENDED-RELEASE TABLETS

### Drug Facts

**Active Ingredient**  
(in each extended-release tablet)  
Guaifenesin 600 mg

Expectorant

**Uses** helps loosen mucus (mucus) and thin bronchial secretions to let 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 mucus (mucous)

#### 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.

**⚠️ pregnant or breast-feeding, ask a health professional before use.**

### Drug Facts (continued)

**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
- this product can be administered without regard for the timing of meals
- adults and children 12 years of age and over: 1 or 2 tablets every 12 hours, Do not exceed 4 tablets in 24 hours.
- children under 12 years of age: do not use

#### Other information

- tamper evident: do not use if seal on bottle printed "SEALED FOR YOUR PROTECTION" is broken or missing.
- store between 20 to 25 °C (68 to 77 °F)

#### Inactive ingredients

carboxymethylcellulose, hydroxypropylmethylcellulose, polyvinylpyrrolidone, croscarmellose, polydioxanone

#### Questions?

1-800-860-2600  
Hours: 8am - 4pm, EST  
You may also report side effects to this phone number.

\*This product is not manufactured or distributed by Reckitt Benckiser, the distributor of Mucinex® Extended Release 600 mg Tablets.

Distributed by: Guardian Drug Company,  
2 Charles Ct, Dayton, NJ 08810 USA

LOT:

BLANK AREA  
FOR LOT/EXP

EXP:

MFR# 53041 REV 0919

# MUCUS RELIEF 600 MG

quaifenesin tablet, extended release

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:53041-190
<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	600 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE</b> (UNII: 0A5MM307FC)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POVIDONE K90</b> (UNII: RDH86HJV5Z)	

Product Characteristics					
Color		WHITE	Score		no score
Shape		CAPSULE	Size		17mm
Flavor			Imprint Code		G233
Contains					
Packaging					
#	Item Code	Package Description		Marketing Start Date	Marketing End Date
1	NDC:53041-190-32	2 in 1 CARTON		06/24/2020	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product			
2	NDC:53041-190-38	4 in 1 CARTON		06/24/2020	
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product			
3	NDC:53041-190-16	100 in 1 BOTTLE; Type 0: Not a Combination Product		06/24/2020	
4	NDC:53041-190-20	500 in 1 BOTTLE; Type 0: Not a Combination Product		06/24/2020	
Marketing Information					
Marketing Category		Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
ANDA		ANDA209215		06/24/2020	

MUCUS RELIEF MAXIMUM STRENGTH 1200 MG			
guaifenesin tablet, extended release			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53041-191
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)		GUAIFENESIN	1200 mg
Inactive Ingredients			
Ingredient Name			Strength
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			



**POVIDONE K90** (UNII: RDH86HJV5Z)

### Product Characteristics

<b>Color</b>	WHITE	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	22mm
<b>Flavor</b>		<b>Imprint Code</b>	G234
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53041-191-58	2 in 1 CARTON	06/24/2020	
1		7 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:53041-191-48	4 in 1 CARTON	06/24/2020	
2		7 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:53041-191-37	28 in 1 BOTTLE; Type 0: Not a Combination Product	04/30/2024	
4	NDC:53041-191-62	70 in 1 BOTTLE; Type 0: Not a Combination Product	04/30/2024	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA209215	06/24/2020	

**Labeler** - Guardian Drug Company (119210276)

### Establishment

Name	Address	ID/FEI	Business Operations
Guardian Drug Company		119210276	MANUFACTURE(53041-190, 53041-191)

Revised: 4/2024

Guardian Drug Company