

**GUAIFENESIN EXTENDED RELEASE- guaifenesin tablet, extended release
PUBLIX SUPER MARKETS, INC**

GUAIFENESIN EXTENDED-RELEASE TABLETS

Drug Facts

Active ingredient (in each extended-release tablet)

Guaifenesin, USP 1200 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 extended-release 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 extended-release tablet every 12

hours. Do not exceed 2 extended-release tablets in 24 hours.

- children under 12 years of age: do not use

Other information

- store between 68-77°F (20-25°C)
- **TAMPER EVIDENT: DO NOT USE IF CARTON IS OPEN OR IF PRINTED SEAL ON BLISTER IS BROKEN OR MISSING.**

Inactive ingredients

colloidal silicon dioxide, FD&C blue #2 aluminum lake, hypromellose, magnesium stearate, povidone.

**DISTRIBUTED BY PUBLIX SUPER MARKETS, INC.,
3300 PUBLIX CORPORATE PARKWAY
LAKELAND, FL 33811**

PRINCIPAL DISPLAY PANEL - 1200 mg Extended-Release Tablet Blister Pack Carton

NDC 41415-997-14

12-HOUR MAXIMUM STRENGTH

mucusrelief

GUAIFENESIN EXTENDED-RELEASE

TABLETS, 1200 mg

EXPECTORANT

12 Hour

- Relieves Chest Congestion
- Thins and Loosens Mucus

14

EXTENDED-RELEASE

TABLETS

†Compare to the active ingredient of Mucinex[®] Maximum Strength



12-HOUR MAXIMUM STRENGTH
mucusrelief
 GUAIFENESIN EXTENDED-RELEASE TABLETS, 1200 mg
 EXPECTORANT

5261031



12-HOUR MAXIMUM STRENGTH
mucusrelief
 GUAIFENESIN EXTENDED-RELEASE TABLETS, 1200 mg
 EXPECTORANT



NDC 41415-997-14

12-HOUR MAXIMUM STRENGTH
mucusrelief
 GUAIFENESIN EXTENDED-RELEASE
 TABLETS, 1200 mg
 EXPECTORANT



12-HOUR MAXIMUM STRENGTH
mucusrelief
 GUAIFENESIN EXTENDED-RELEASE TABLETS, 1200 mg
 EXPECTORANT



ACTUAL SIZE

12 Hour

- Relieves Chest Congestion
- Thins and Loosens Mucus

14 EXTENDED-RELEASE
 TABLETS

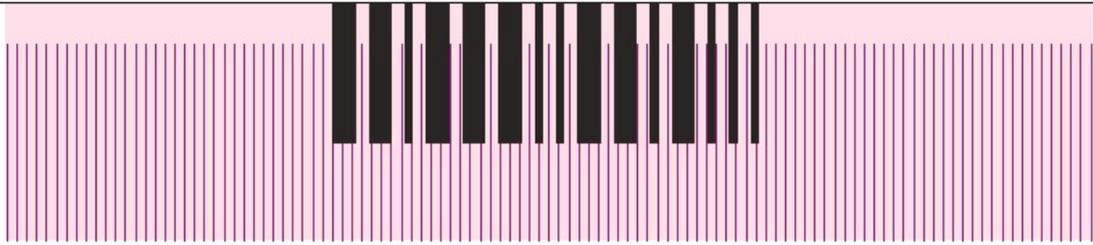
¹Compare to the active
 ingredient of Mucinex®
 Maximum Strength

14 EXTENDED-RELEASE
 TABLETS

14 EXTENDED-RELEASE
 TABLETS

NVZ





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Guaifenesin, USP 1200 mg.....	Expectorant

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Keep the carton. It contains important information. See end panel for expiration date.

†All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Mucinex®.

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LAKE LAND, FL 33811
1-888-267-3037 publix.com

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PRODUCT OF INDIA

CODE No.: PB/DRUGS/1809-OSP





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GUAIFENESIN EXTENDED RELEASE

guaifenesin tablet, extended release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41415-997
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
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	

Product Characteristics

Color	white	Score	no score
Shape	OVAL	Size	22mm
Flavor		Imprint Code	RH99
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC 41415			

1	NDC:41415-997-14	1 in 1 CARTON	12/10/2017	
1		14 in 1 BLISTER PACK; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA209254		12/10/2017	

Labeler - PUBLIX SUPER MARKETS, INC (006922009)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(41415-997)

Revised: 5/2025

PUBLIX SUPER MARKETS, INC