# MUCUS RELIEF- guaifenesin tablet, film coated Chain Drug Consortium

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**Premier Value 44-532** 

## Active ingredient (in each immediate-release tablet)

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

## Ask a doctor before use if you have

- cough accompanied by too much phlegm (mucus)
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema

# Stop use and ask a doctor if

cough persists more than 7 days, tends to recur, or is accompanied by a 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.

#### **Directions**

- take with a full glass of water
- adults and children 12 years and over: 1 tablet every 4 hours. Do not take more than 6 tablets in 24 hours.
- children under 12 years: do not use

#### Other information

store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

see end flap for expiration date and lot number

## Inactive ingredients

FD&C blue #1 aluminum lake, hypromellose, magnesium stearate, maltodextrin, microcrystalline cellulose, polyethylene glycol, povidone, silicon dioxide, sodium starch glycolate, stearic acid

#### Questions or comments?

1-800-426-9391

#### Principal Display Panel

Premier Value<sup>®</sup>

Immediate release

Mucus Relief

### **GUAIFENESIN, 400 mg**

Expectorant

Relieves Chest Congestion Thins and Loosens Mucus

actual size **50** Tablets

#### PV

INDEPENDENTLY TESTED SATISFACTION GUARANTEED

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

50844 ORG011853215 Distributed By: Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 19087

If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.



**Premier Value 44-532** 

# MUCUS RELIEF guaifenesin tablet, film coated Product Information Product Type HUMAN OTC DRUG Item Code (Source) 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
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics				
Color	blue	Score	2 pieces	
Shape	ROUND	Size	13mm	
Flavor		Imprint Code	44;532	
Contains				

l	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:68016- 545-50	1 in 1 CARTON	01/09/2023	
	L	50 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 Drug	M012	01/09/2023			

# Labeler - Chain Drug Consortium (101668460)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		038154464	pack(68016-545)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations

manufacture(68016-545), p	oack(68016-545)
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LNK	International.	Inc
LINK	international.	IIIC.

832867837

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		832867894	manufacture(68016-545)

Establishment			
Name	Address	ID/FEI	Business Operations
LNK International, Inc.		967626305	pack(68016-545)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
LNK International, Inc.		117025878	manufacture(68016-545)

Revised: 1/2024 Chain Drug Consortium