

**MAXIMUM STRENGTH MUCUS RELIEF- guaifenesin tablet, extended release**  
**CHAIN DRUG MARKETING ASSOCIATION INC**

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**1204A-QCH-2021-1108**

***Drug Facts***

**Active ingredient (in each extended-release tablet)**

Guaifenesin 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 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**

■ store between 20-25°C (68-77°F)

■ retain carton for complete product information and warnings

### **Inactive ingredients**

carbomer homopolymer type B, hypromellose, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

### **PRINCIPAL DISPLAY PANEL**

QUALITY CHOICE®

NDC 63868-148-14

†Compare to the Active Ingredient in MUCINEX® Maximum Strength

12 Hour

Maximum Strength

Mucus Relief

Expectorant

Guaifenesin Extended-Release Tablets, 1200 mg

- Relieves Chest Congestion
- Thin & Loosens Mucus
- Immediate and Extended Release

Actual Size

14 EXTENDED-RELEASE TABLETS



## MAXIMUM STRENGTH MUCUS RELIEF

guaifenesin tablet, extended release

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63868-148
<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	1200 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>CARBOMER HOMOPOLYMER TYPE B (ALLYL SUCROSE CROSSLINKED)</b> (UNII: Z135WT9208)	
<b>HYPROMELLOSE 2910 (15 MPA.S)</b> (UNII: 365FW2JZ0W)	
<b>SODIUM STARCH GLYCOLATE TYPE A</b> (UNII: H8AV0SQX4D)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	OVAL (Elliptical)	<b>Size</b>	22mm
<b>Flavor</b>		<b>Imprint Code</b>	G;1200
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-148-14	2 in 1 CARTON	09/17/2021	
1		7 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	ANDA213420	09/17/2021	

**Labeler** - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

Revised: 11/2021

CHAIN DRUG MARKETING ASSOCIATION INC