

GUAIFENESIN- guaifenesin liquid
Method Pharmaceuticals, LLC

Guaifenesin

NDC 58657-509-16

Guaifenesin

Liquid USP

100 mg/5 mL

Expectorant

Sugar Free • Alcohol Free

Cherry Flavor

Loosens and Relieves Chest Congestion

16 fl. oz. (473 mL)

Drug Facts

Active ingredient (in each 5 mL)

Guaifenesin 100 mg

Purpose

Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive.

Warnings

Ask a doctor before use if you have

- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis, or emphysema

Stop use and ask a doctor if

- cough lasts more than 7 days, comes back, or is accompanied by 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

- do not take more than 6 doses in any 24-hour period
- age dose adults and children 2 to 4 teaspoonfuls
- 12 years and over every 4 hours
- children 6 years to 1 to 2 teaspoonfuls
- under 12 years every 4 hours
- children 2 years to ½ to 1 teaspoonful
- under 6 years every 4 hours
- children under 2 years ask a doctor

Drug Facts(continued)

Other information

- store at 20°-25°C (68°-77°F)
- packaged with tamper evident seal under cap

Inactive ingredients

Bitter Mask, Cherry Flavor, Citric Acid, FD&C Red#40, Glycerin, Propylene Glycol, Purified Water, Sodium Benzoate, Sodium Citrate, Sodium Saccharin, Sorbitol Solution

Questions or Comments

1-877-250-3427

Manufactured For:

Method Pharmaceuticals, LLC

Fort Worth, Texas 76118 Rev. 09/16

PRINCIPAL DISPLAY PANEL

NDC 58657- 509- 16

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Directions

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age	dose
adults and children 12 years and over	2 to 4 teaspoonsful every 4 hours
children 6 years to under 12 years	1 to 2 teaspoonsful every 4 hours
children 2 years to under 6 years	½ to 1 teaspoonful every 4 hours
children under 2 years	ask a doctor



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Lot:
Exp.:

GUAIFENESIN

guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58657-509
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	100 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	
SORBITOL (UNII: 506T60A25R)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

Product Characteristics

Color	red	Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58657-509-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	02/09/2016	

Labeler - Method Pharmaceuticals, LLC (060216698)

Revised: 10/2025

Method Pharmaceuticals, LLC