TUSNEL EX- guaifenesin liquid
A-S Medication Solutions

In each 5 mL

Guaifenesin - 100 mg

Purpose

Expectorant

Uses

- temporarily relieves cough due to minor thraot and bronchial irritations as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive

Warnings

Do not use if you are now taking a prescritpion monamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's diesease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescirption drug contains an MAOI, aska doctor or pharmacist before taking this product.

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 for more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. A persistent cough may be a sign of a serious condition.

If pregnant or breast-feeing, 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 exceed more than 6 doses in any 24-hour period.

Age	Dose
Adults and children over 12 years and over	2 teaspoonfuls (10 mL) every 4 hours
Children under 12 years	Ask a doctor

Inactive ingredients anhydrous citric acid, flavor, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate digydrate, and a sucralose.

Questions or comments? 1-8660595-5598

HOW SUPPLIED

Product: 50090-6542

NDC: 50090-6542-0 118 mL in a BOTTLE

guaifenesin liquid

TUSNEL EX

guaifenesin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50090-6542(NDC:54859-507)
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**
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090-6542-0	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/05/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	05/01/2019	

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-6542)

Revised: 12/2023 A-S Medication Solutions