

TUSNEL EX- guaifenesin liquid
Llorens Pharmaceutical International Division, Inc.

Llorens-TusnelEX 507

In each 5 mL

Guaifenesin - 100 mg

Purpose

Expectorant

Uses

- temporarily relieves cough due to minor throat 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 prescription monamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a 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-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 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

NDC: 54859-507-04



Drug Facts	
Active Ingredient (per 5 mL teaspoonful)	Purpose
Guaifenesin, USP 100 mg	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 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-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 (1-800-222-1222) right away.	

Drug Facts (continued)	
Directions	
• Do not take more than 6 doses in any 24-hour period.	
• Adults and children 12 years of age and over, take two teaspoonfuls (10 mL) every 4 hours or as directed by a doctor.	
• Children under 12 years of age, do not use unless directed by a doctor.	
Other Information	
• store at room temperature 20-25°C (68-77°F)	
• Tamper-Evident by seal under cap. Do not use if there is evidence of tampering.	
Inactive Ingredients: artificial and natural apple banana flavor, citric acid anhydrous, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate dihydrate, and sucralose.	
Questions or comments: 1-866-595-5598	

TUSNEL[®] EX
 Guaifenesin, USP
 100 mg / 5 mL
 Sugar Free

Item #: L-153 REV. 07/24

Manufactured by:
 Llorens Pharmaceutical International Division, Inc.
 7080 NW 37th Ct, Miami, FL 33147

Expectorant



Apple Banana Flavor
 4 oz (118 mL)



TUSNEL EX

guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	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:54859-507-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2019	

Marketing Information

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

Labeler - Llorens Pharmaceutical International Division, Inc. (037342305)

Revised: 12/2025

Llorens Pharmaceutical International Division, Inc.