



**Keep out of reach of children.** In case of accidental overdose, seek advice or a doctor or contact a Poison Control Center right away.

**If pregnant or breast-feeding,** ask a doctor before use.

**Directions** Do not exceed 6 doses in 24 hours.

adults and children over 12 years of age take 1 teaspoonful (5 mL) every 6-8 hours  
 children 6 to under 12 years of age take 1/2 teaspoonful (2.5 mL) every 6-8 hours  
 children under 6 years of age consult a doctor before use

**Inactive Ingredients** citric acid, FDandC yellow No 6, glycerin, menthol, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate and sucralose

**Questions or comments?** 1-305-403-3788

Monday - Friday 9:00AM - 5:00 PM EST.

**Manufactured For:** Advanced Generic Corporation, Miami, FL 33147

www.advancedgeneric.com

NDC 45737-242-16

**Drug Facts**

**Active ingredients (in each 5 mL teaspoonful)**

Dextromethorphan HBR, USP.....	28 mg.....	Cough Suppressant
Guaifenesin, USP.....	388 mg.....	Expectorant
Phenylephrine HCl, USP.....	10 mg.....	Nasal Decongestant

**Uses** • temporary relieves cough due to minor throat and bronchial irritations as may occur with the common cold or inhaled irritants • helps loosen phlegm (mucus) and thin bronchial secretions to make cough more productive • temporarily relieves nasal congestion due to the common cold

**Warnings** • Do not use if you are now taking a monoamine 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** • heart disease • high blood pressure • thyroid disease • diabetes • difficulty in urination due to enlargement of prostate gland • a cough with too much phlegm (mucus) • a persistent or chronic cough as occurs with smoking, asthma, chronic bronchitis, or emphysema

**Ask a doctor before use if you are** taking sedatives, tranquilizers or drugs for depression or MAOI drugs.

# BioGruss

## NF

- Cough Suppressant
- Expectorant
- Nasal Decongestant

**Alcohol FREE • Sugar FREE**

Contains the same active ingredients  
Giltuss® New Improved Formula<sup>†</sup>

Grape Flavor

Manufactured For:  
 advanced generic corporation  
 Miami, FL 33147  
 www.advancedgeneric.com

16 fl. oz. (473 mL)

**Drug Facts** (continued)

**Stop use and ask a doctor if** • nervousness, dizziness, or sleeplessness occur • cough lasts for more than 7 days, comes back or occurs with a fever, rash or headache that lasts. These could be signs of a serious condition.

**If pregnant or breast-feeding,** ask a doctor before use  
**Keep out of the reach of children.** In case of accidental overdose, seek advice of a doctor or contact a Poison Control Center right away.

**Directions** Do not exceed 4 doses in 24 hours.

adults and children over 12 years of age	take 1 teaspoonful (5 mL) every 6-8 hours
children 6 to under 12 years of age	take 1/2 teaspoonful (2.5 mL) every 6-8 hours
children under 6 years of age	consult a doctor before use

**Other information** • store at controlled room temperature 15°-30° C (59°- 86° F).

**Tamper Evident Feature:** Do not use if seal under cap (or cello-band over cap) is torn, broken or missing.

**Pharmacist-** Preserve and dispense in tight-light resistant container with a child resistant cap as defined in the USP.

**Inactive ingredients** Citric acid, FD&C yellow #6, glycerin, L-menthol, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, and sucralose.

**Questions or comments?** 1-305-403-3788  
 Monday - Friday 9:00 am - 5:00 pm EST.

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THIS IS A BULK CONTAINER NOT INTENDED FOR DISPENSING

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<b>BIOGTUSS NF</b>			
dextromethorphan, guaifenesin, phenylephrine liquid			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:45737-242
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	28 mg in 5 mL
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	388 mg in 5 mL
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 5 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>MENTHOL</b> (UNII: L7T10EIP3A)	
<b>METHYLPARABEN</b> (UNII: A2I8C7HI9T)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>PROPYLPARABEN</b> (UNII: Z8IX2SC1OH)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	

**Product Characteristics**

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	GRAPE (Grape Flavor)	<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:45737-242-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2010	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/01/2010	

**Labeler** - Advanced Generic Corporation (831762971)