**BIOGTUSS NF- dextromethorphan, guaifenesin, phenylephrine liquid**
**Advanced Generic Corporation**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**Active Ingredients: (in each 5 mL tsp.)** **Purpose**

Dextromethorphan HBr 28 mg..... Cough Suppressant

Guaifenesin 388 mg ..... Expectorant

Phenylephrine HCl 10 mg.....Nasal Decongestant

**Purpose**

Cough suppressant

Expectorant

Nasal Decongestant

**Uses**

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

**Warnings**

**Do not use** If you are now taking a prescription 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 you are taking a prescription drug that contains an MAOI; ask your 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.

**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

**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 |

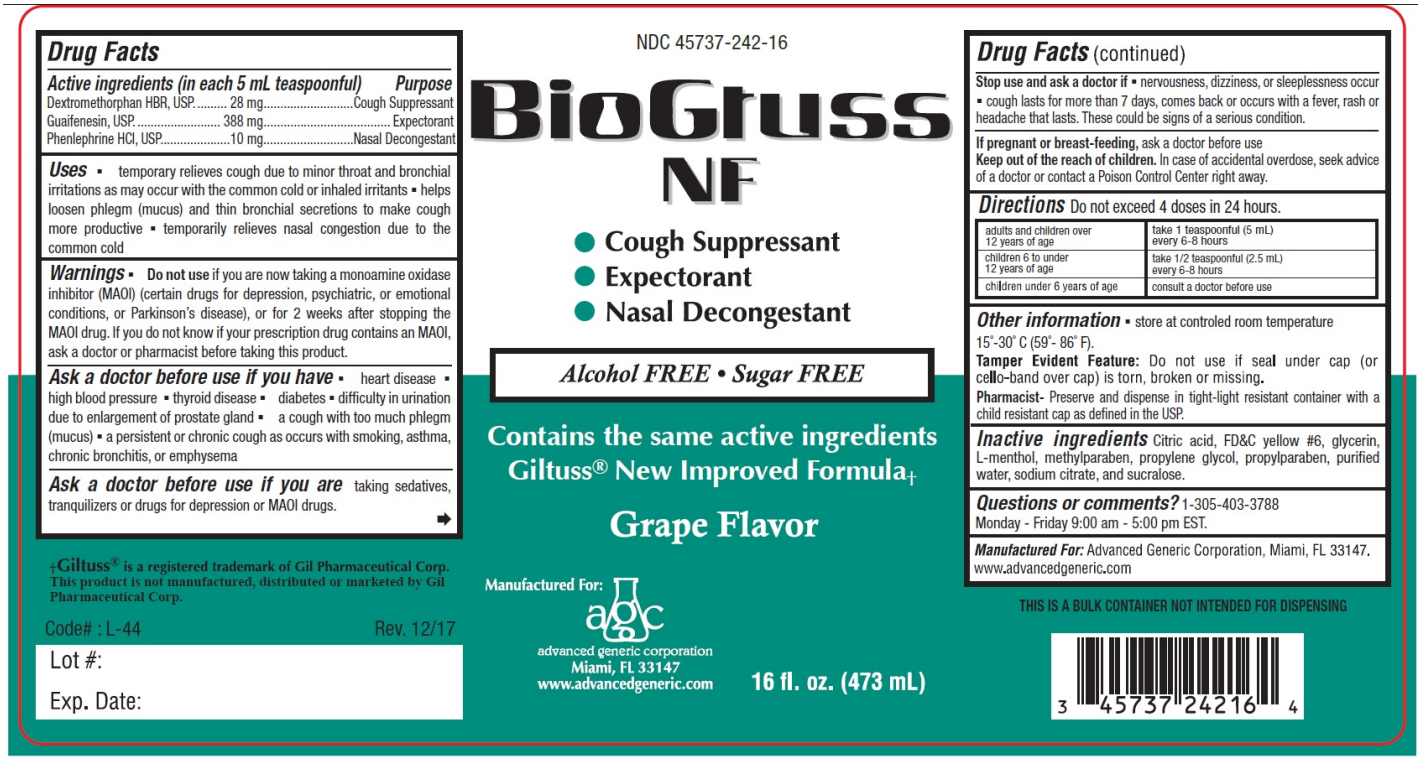
**Inactive Ingredients** citric acid, FDandC yellow No 6, glycerin, menthol, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate and sucralose

**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




# BIOGTUSS NF

dextromethorphan, guaifenesin, phenylephrine liquid

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:45737-242 |
| **Route of Administration** | ORAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **Dextromethorphan Hydrobromide** (UNII: 9D2RTI9KYH) (Dextromethorphan - UNII:7355X3ROTS) | Dextromethorphan Hydrobromide | 28 mg in 5 mL |
| **Guaifenesin** (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ) | Guaifenesin | 388 mg in 5 mL |
| **Phenylephrine Hydrochloride** (UNII: 04JA59TNSJ) (Phenylephrine - UNII:1WS297W6MV) | Phenylephrine Hydrochloride | 10 mg in 5 mL |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |
| **FD&C YELLOW NO. 6** (UNII: H77VEI93A8) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **MENTHOL** (UNII: L7T10EIP3A) | |
| **METHYLPARABEN** (UNII: A2I8C7HI9T) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **PROPYLPARABEN** (UNII: Z8IX2SC1OH) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **SODIUM CITRATE** (UNII: 1Q73Q2JULR) | |
| **SUCRALOSE** (UNII: 96K6UQ3ZD4) | |

| Product Characteristics | | | |
|---|---|---|---|
| **Color** | | **Score** | |
| **Shape** | | **Size** | |
| **Flavor** | GRAPE (Grape Flavor) | **Imprint Code** | |
| **Contains** | | | |

| 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 final | part341 | 09/01/2010 | |

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

Revised: 12/2020 Advanced Generic Corporation