# BIODESP DM- 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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# **Drug Facts**

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

Dextromethorphan Hydrobromide 15 mg...... Cough Suppressant

Guaifenesin 100 mg ...... Expectorant

Phenylephrine HCl 5 mg...... Decongestant

### **Purpose**

**©Cough Suppressant** 

**Expectorant** 

Nasal Decongestant

#### Uses

- temporarily relieves these symptoms due to the common cold. hay fever (allergic rhinitis), or other upper respiratory allergies;
- cough due to minor throat and bronchial irritation
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bonchial tubes and make coughs more productive
- nasal congestion
- reduces swelling of nasal passages

# **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.

**■When using this product** do not exceed recommended dosage

#### **Ask a doctore before use if you have**

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- trouble urinating due to enlarged prostate gland
- a cough with too much phlegm (mucus)
- a persistent or chronic cough as occurs with smoking, asthma, chronic bronchitis or emphysema.

#### Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occurs
- 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 overdose, get medical help or contact a Poison Control Center right away.

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

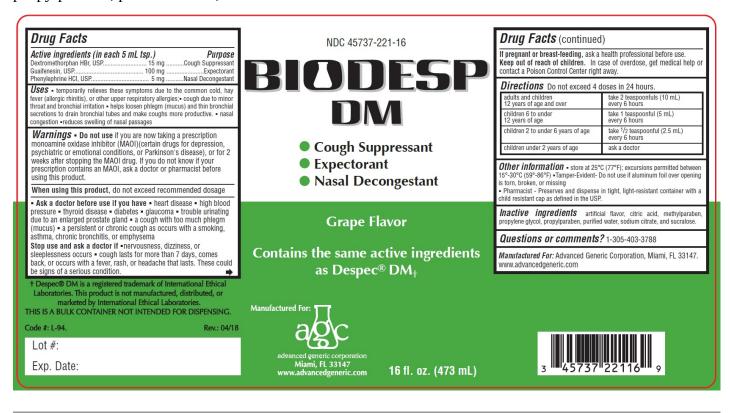
**Directions** Do not exceed 4 doses in 24 hours.

adults and children 12 years of age and over children 6 to under 12 years of age children 2 to under 6 years of age children under 2 years of age take 2 teaspoonfuls (10 mL) every 6 hours take 1 teaspoonful (5 mL) every 6 hours take 1/2 teaspoonful (2.5 mL) every 6 hours ask a doctor

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

Manufactured For: Advanced Generic Corporation, Miami, FL 33147. www.advancedgeneric.com

**Inactive ingredients** artificial flavor, citric acid, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate and sucralose.



# BIODESP DM

dextromethorphan, guaifenesin, phenylephrine liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:45737-221
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingre	dient Name		Basis of Strength	Strength

<b>Dextromethorphan Hydrobromide</b> (UNII: 9D2RTI9KYH) (Dextromethorphan - UNII:7355X3ROTS)	Dextromethorphan Hydrobromide	15 mg in 5 mL
Guaifenesin (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ)	Guaifenesin	100 mg in 5 mL
Phenylephrine Hydrochloride (UNII: 04JA59TNSJ) (Phenylephrine - UNII:1WS297W6MV)	Phenylephrine Hydrochloride	5 mg in 5 mL

Inactive Ingredients			
Ingredient Name	Strength		
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)			
Water (UNII: 059QF0KO0R)			
Methylparaben (UNII: A2I8C7HI9T)			
Propylparaben (UNII: Z8IX2SC1OH)			
Propylene Glycol (UNII: 6DC9Q167V3)			
Sodium Citrate (UNII: 1Q73Q2JULR)			
SUCRALOSE (UNII: 96K6UQ3ZD4)			

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

	Packaging			
ı	# Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date
ı	1 NDC:45737-221-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	0 1/0 1/20 0 9	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	0 1/0 1/20 0 9	

# Labeler - Advanced Generic Corporation (831762971)

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