

**BIODESP DM- dextromethorphan, guaifenesin, phenylephrine liquid**  
**Advanced Generic Corporation**

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**AGC-BiodespDM 221**

**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 bronchial 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 doctor 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**

<b>BIODESP DM</b> dextromethorphan, guaifenesin, phenylephrine liquid
<b>Product Information</b>

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:45737-221	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
DEXTROMETHORPHAN HYDROBROMIDE (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	
ANHYDROUS 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				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:45737-221-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2009	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012		01/01/2009	

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