BIONATUSS DXP- dexbrompheniramine maleate, dextromethorphan, phenylephrine liquid
Advanced Generic Corporation

AGC-BionatussDXP 210

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

Dexbrompheniramine Maleate.....2mg.....Antihistamine

Dextromethorphan HBr.....20mg.....Cough Suppressant

Phenylephrine HCL.....10mg..... Nasal Descongestant

Purpose

Antihistamine

Cough Suppressant

Nasal Decongestant

Uses

- helps to control the relex that causes coughing
- temporarily relieves nasal congestion due to common cold, hay fever, or other upper respiratory allergies (allergic rhinitis)
- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
- runny nose
- itchy, watery eyes
- sneezing
- itching of the nose or throat

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 your prescription contains an MAOI, ask a doctor or pharmacist before using the product.

Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- trouble urinating due to an enlarged 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 docotor before use if you are taking sedatives or tranquilizers

When using this product

- do not exceed recommended dose
- excitability may occur, especially in children
- drowsiness may occur
- avoid alcoholic beverages
- alcohol, sedatives and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating

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.

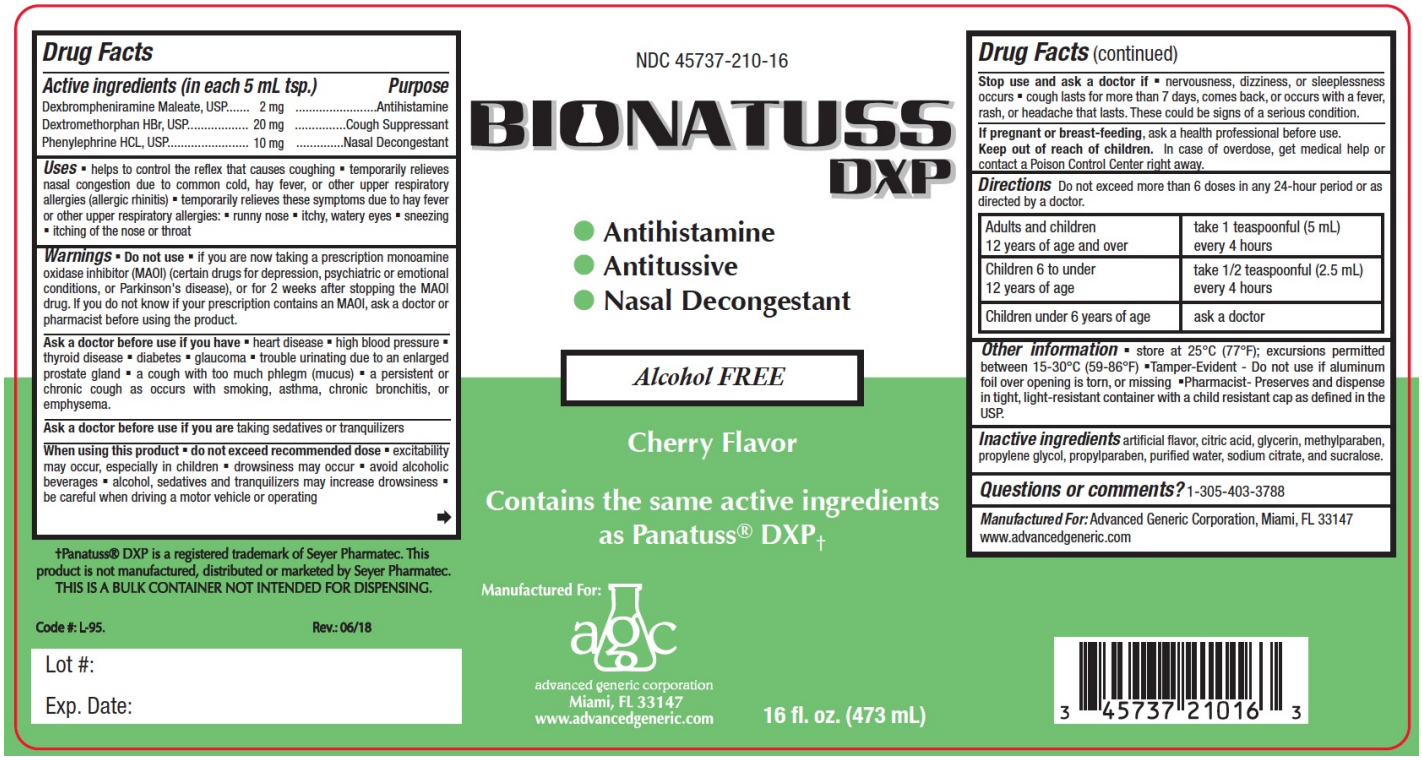
If pregnant or breast-feeding,ask a health professional before use.

Directions: Do not exceed more than 6 doses in any 24-hour period or as directed by a doctor.

Adults and children 12 years of age and over	take 1 teaspoonful (5 mL) every 4 hours
Children 6 to under 12 years of age	take 1/2 teaspoonful (2.5 mL) every 4 hours
Children under 6 years of age	ask a doctor

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

Questions or comments?1-305-403-3788



BIONATUSS DXP

dexbrompheniramine maleate, dextromethorphan, phenylephrine liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:45737-210
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXBROMPHENIRAMINE MALEATE (UNII: BPA9UT29BS) (DEXBROMPHENIRAMINE - UNII:75T64B71RP)	DEXBROMPHENIRAMINE MALEATE	2 mg in 5 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
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	RASPBERRY (Flavor)	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:45737-210-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/01/2009	

Labeler - Advanced Generic Corporation (831762971)

Revised: 12/2024 Advanced Generic Corporation