BRANTUSSIN DM- dexbrompheniramine maleate, dextromethorphan hbr, phenylephrine hcl liquid
Brandywine Pharmaceuticals, LLC

BranTussin DM

Active Ingredient

Dexbrompheniramine Maleate 2 mg
Dextromethorphan Hydrobromide 15 mg
Phenylephrine Hydrochloride 7.5 mg

Uses

temporarily relieves these symptoms due to the common cold, hayfever (allergic rhinitis) or other upper respiratory allergies:

- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes
- cough due to minor throat and
- bronchial irritation
- nasal congestion
- reduces swelling of nasal passages

Purposes

Antihistamine

Cough Suppressant

Nasal Decongestant

Warnings

Do not exceed recommended dosage.

Do not use this product

if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional condition, 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

- a breathing problem such as emphysema or chronic bronchitis
- glaucoma

- trouble urinating due to an enlarged prostate gland
- a cough that lasts or is chronic such as occurs with smoking, asthma or emphysema
- a cough that occurs with too much phlegm (mucus)
- heart disease
- high blood pressure
- thyroid disease
- diabetes mellitus

Stop use and ask a doctor if

 nervousness, dizziness, or sleeplessness occur

 cough or nasal congestion persists for more than 1 week, tends to recur, or is accompanied by a fever, rash or persistent headache. A persistent cough may be a sign of a serious condition.

 new symptoms occur

Keep this and all drugs out of the reach of children

In case of accidental overdose, get medical help or contact a Poison Control Center immediately.

When using this product

 excitability may occur, especially in children

 may cause marked drowsiness

 avoid alcoholic drinks
 alcohol, sedatives, and tranquilizers may increase the drowsiness effect

 be careful when driving a motor vehicle or operating machinery

Ask a doctor or pharmacist before use

if you are taking sedatives or tranquilizers.

If pregnant or breastfeeding

ask a health professional before use.

Directions

Do not exceed recommended dosage

Adults and children 12 years of age and over	1 teaspoonful (5 mL) every 4 to 6 hours, not to exceed 6 teaspoonfuls in 24
Children 6 to under 12 years of age	1/2 teaspoonful (2.5 mL) every 4 to 6 hours, not to exceed 3 teaspoonfuls in 24 hours
Children under 6 years of age	Consult a doctor.

Other Information

Store at 59° - 86°F (15° - 30°C)

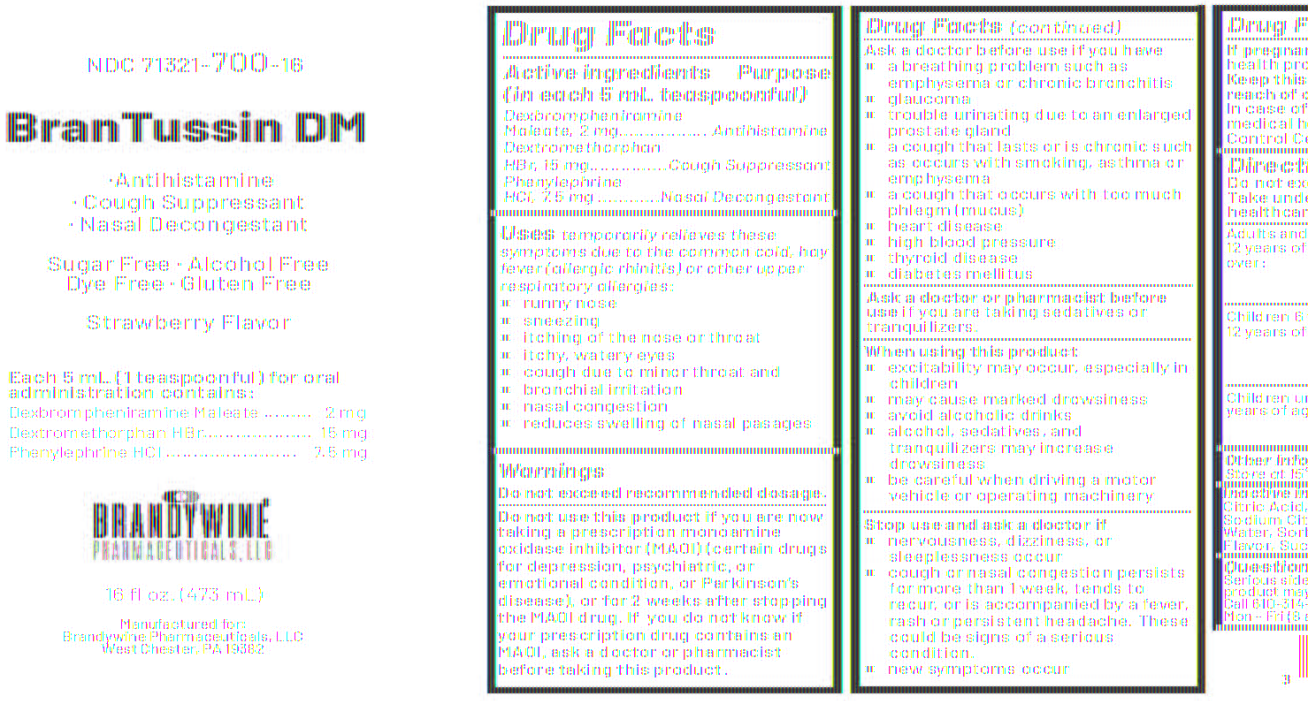
Inactive Ingredients

Anhydrous Citric Acid, Glycerin, Flavor, Propylene Glycol, Purified Water, Sodium Benzoate, Sodium Citrate Dihydrate, Sodium Saccharin, Sorbitol Solution.

Questions? Comments?

Serious side effects associated with use of this product may be reported to this number. Call 1-800-882-1041 Mon. - Fri. (8 a.m. to 5 p.m. CST).

Package Label



BRANTUSSIN DM

dexbrompheniramine maleate, dextromethorphan hbr, phenylephrine hcl liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:71321-700
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	7.5 mg in 5 mL

DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg in 5 mL
DEXBROMPHENIRAMINE MALEATE (UNII: BPA9UT29BS) (DEXBROMPHENIRAMINE - UNII:75T64B71RP)	DEXBROMPHENIRAMINE MALEATE	2 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SORBITOL (UNII: 506T60A25R)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71321-700-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/14/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/14/2022	

Labeler - Brandywine Pharmaceuticals, LLC (080581956)

Revised: 1/2024 Brandywine Pharmaceuticals, LLC