

**DURATUSS DM- dextbrompheniramine maleate, dextromethorphan hydrobromide and pseudoephedrine hydrochloride liquid**  
**Poly Pharmaceuticals, Inc.**

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**DURATUSS DM**

**Active Ingredients**

**(in each 5 mL teaspoonful)**

Dextbrompheniramine Maleate 1 mg

Dextromethorphan Hydrobromide 10 mg

Pseudoephedrine Hydrochloride 30 mg

**Purpose**

Antihistamine  
Antitussive  
Nasal Decongestant

**Uses**

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

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

**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 conditions, 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

**Ask a doctor or pharmacist before use****if you are**

taking sedatives or tranquilizers.

**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

**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

**If pregnant or breast-feeding,**

ask a health professional before use.

**Keep out of reach of children.**

In case of overdose, get medical help or contact a Poison Control Center right away.

**Directions****Do not exceed recommended dosage.**

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Adults and	2 teaspoonful (10 mL)
children 12	every 4 to 6 hours,
years of age	not to exceed 8
	teaspoonfuls in a 24

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

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### **Other information**

Store at 59° - 86°F (15° - 30°C)

### **Inactive ingredients**

Citric acid, flavors, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose.

### **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).

### **PRINCIPAL DISPLAY PANEL**

**NDC 50991-142-16**

**Duratuss DM**

**LIQUID**

**Antihistamine • Antitussive**

**Nasal Decongestant**

**Raspberry Bubble Gum Flavor**

**SUGAR FREE / ALCOHOL FREE**

**DYE FREE / GLUTEN FREE**

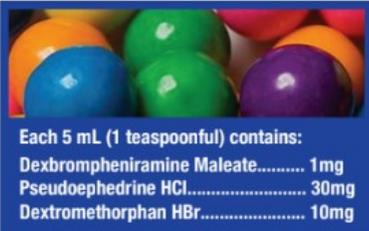
**16 fl oz. (473 mL)**

Tamper evident by foil seal under cap. Do not use if foil seal is broken or missing. Dispense in a tight, light-resistant container with a child-resistant cap.  
**THIS BOTTLE IS NOT TO BE DISPENSED TO CONSUMER.**  
**Distributed by: Poly Pharmaceuticals, Owens Cross Roads, AL 35763**

NDC 50991-142-16

## DURATUSS DM

Antihistamine • Nasal Decongestant • Antitussive



Each 5 mL (1 teaspoonful) contains:  
Dexbrompheniramine Maleate..... 1mg  
Pseudoephedrine HCl..... 30mg  
Dextromethorphan HBr..... 10mg



Raspberry Bubblegum Flavor

**Alcohol Free • Dye Free  
Sugar Free • Gluten Free**

Tamper evident by foil seal under cap. Do not use if foil seal is broken or missing. Dispense in a tight, light-resistant container with a child-resistant cap. This bottle is not to be dispensed to consumer.

Distributed by:  
Poly Pharmaceuticals  
Huntsville, AL 35763



16 fl oz. (473 mL)

### Drug Facts

Active ingredients (in each 5 mL teaspoonful)	Purpose
Dexbrompheniramine Maleate..... 1 mg	
Pseudoephedrine HCl..... 30 mg	
Dextromethorphan HBr..... 10 mg	

Dexbrompheniramine Maleate is the Antihistamine, Pseudoephedrine HCl is the Nasal Decongestant and Dextromethorphan HBr is the Antitussive.

**Uses** temporarily relieves these symptoms due to the common cold, hay fever (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

**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 conditions, 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  
■ difficulty in urination due to the enlargement of the 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  
■ heart disease ■ high blood pressure ■ thyroid disease  
■ diabetes

**Ask a doctor or pharmacist before use if you are taking sedatives or tranquilizers.**

**When using this product**  
■ excitability may occur, especially in children  
■ may cause marked drowsiness ■ avoid alcoholic drinks  
■ alcohol, sedatives, and tranquilizers may increase drowsiness effect  
■ be careful when driving a motor vehicle or operating machinery

### Drug Facts (continued)

**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. These could be signs of a serious condition.  
■ new symptoms occur

**If pregnant or breast-feeding, ask a health professional before use.**  
**Keep out of the reach of children.**  
**In case of overdose, get medical help or contact a Poison Control Center right away.**

**Directions**  
**Do not exceed recommended dosage.**

AGE	DOSE
Adults and children 12 years of age and over:	2 teaspoonfuls (10 mL) every 4 to 6 hours, not to exceed 8 teaspoonfuls in 24 hours, or as directed by a doctor.
Children 6 to under 12 years of age:	1 teaspoonful (5 mL) every 4 to 6 hours, not to exceed 4 teaspoonfuls in 24 hours, or as directed by a doctor.
Children under 6 years of age:	Consult a doctor.

**Other information**  
Store at 59° - 86°F (15° - 30°C)

**Inactive ingredients**  
Citric acid, flavors, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose.

**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)

Iss. 07/25

## DURATUSS DM

dexbrompheniramine maleate, dextromethorphan hydrobromide and pseudoephedrine hydrochloride liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:50991-142
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DEXBROMPHENIRAMINE MALEATE</b> (UNII: BPA9UT29BS) (DEXBROMPHENIRAMINE - UNII: 75T64B71RP)	DEXBROMPHENIRAMINE MALEATE	1 mg in 5 mL
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII: 7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 5 mL
<b>PSEUDOEPHEDRINE HYDROCHLORIDE</b> (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII: 7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	30 mg in 5 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>SORBITOL</b> (UNII: 506T60A25R)	

## Product Characteristics

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	BUBBLE GUM (Raspberry Bubblegum)	<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50991-142-20	20 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2025	
2	NDC:50991-142-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2025	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	10/15/2025	

**Labeler** - Poly Pharmaceuticals, Inc. (198449894)

Revised: 10/2025

Poly Pharmaceuticals, Inc.