

**WESTUSSIN DM NF- westussin dm nf liquid**  
**Westminster Pharmaceuticals, LLC**

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**WesTussin DM NF**

**Drug Facts**

<b>Active ingredients (in each 5 mL teaspoonful)</b>	<b>Purpose</b>
Dexbrompheniramine Maleate 2 mg	Antihistamine
Dextromethorphan Hydrobromide 15 mg	Cough Suppressant
Phenylephrine Hydrochloride 7.5 mg	Nasal Decongestant

**Uses**

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
- use caution 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 breastfeeding** 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**

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 hours
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, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose.

### **Questions?**

Call weekdays from 9 AM to 5 PM EST at 1-844-221-7294. You may also report serious side effects to this phone number.

**PRINCIPAL DISPLAY PANEL - 473 mL Bottle Label**

NDC 69367-353-16

WesTussin DM NF

Antihistamine • Cough Suppressant

• Nasal Decongestant

SUGAR FREE / ALCOHOL FREE

DYE FREE / GLUTEN FREE

Each 5 mL (1 teaspoonful) contains:

Dexbrompheniramine Maleate

2 mg

Dextromethorphan HBr

15 mg

Phenylephrine HCl

7.5 mg

Strawberry Flavor

TAMPER EVIDENT: Tamper evident by foil seal under cap. Do not use if foil seal is broken or missing.

16 OZ (473 mL)

Westminster

Pharmaceuticals

NDC 69367-353-16

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Manufactured for:  
**Westminster Pharmaceuticals, LLC**  
 Nashville, TN 37217

Rev. 05/24



## WESTUSSIN DM NF

westussin dm nf liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69367-353
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1W5297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	7.5 mg in 5 mL
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	15 mg in 5 mL
<b>DEXBROMPHENIRAMINE MALEATE</b> (UNII: BPA9UT29BS) (DEXBROMPHENIRAMINE - UNII:75T64B71RP)	DEXBROMPHENIRAMINE MALEATE	2 mg in 5 mL

**Inactive Ingredients**

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>GLYCERIN</b> (UNII: PDC6A3COOX)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KOOR)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	

<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>TRISODIUM CITRATE DIHYDRATE</b> (UNII: B22547B95K)	

### Product Characteristics

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	STRAWBERRY	<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69367-353-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2023	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH DRUG	M012	07/19/2023	

**Labeler** - Westminster Pharmaceuticals, LLC (079516651)

Revised: 8/2024

Westminster Pharmaceuticals, LLC