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

WesTussin DM NF

Drug Facts

Active ingredients (in each 5 mL teaspoonful)	***Purpose***
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

Citric acid anhydrous, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose, trisodium citrate dihydrate.

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

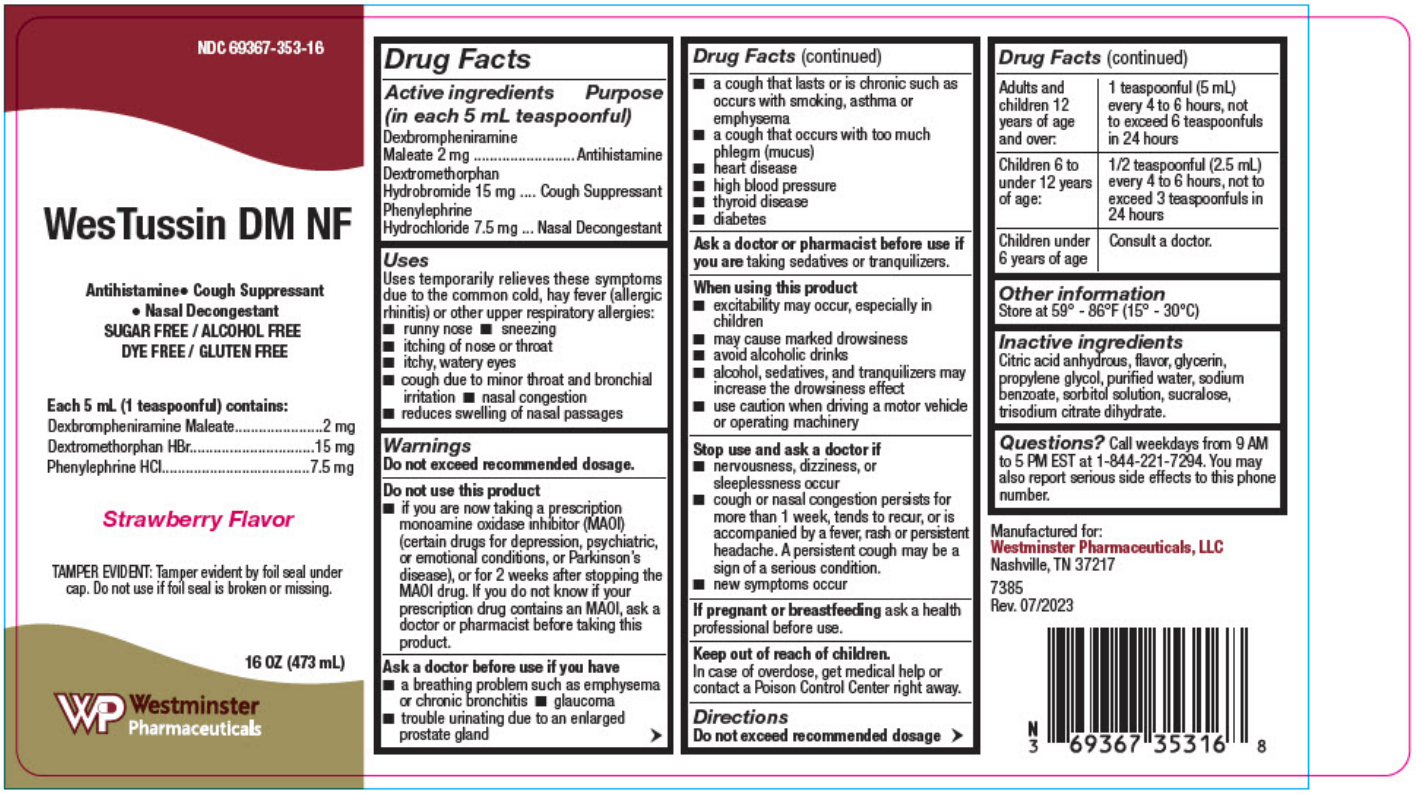
Each 5 mL (1 teaspoonful) contains:
Dexbrompheniramine Maleate
2 mg
Dextromethorphan HBr
15 mg
Phenylephrine HCl
7.5 mg

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



WESTUSSIN DM NF

westussin dm nf liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69367-353
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
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	

Product Characteristics			
Color		**Score**	
Shape		**Size**	
Flavor	STRAWBERRY	**Imprint Code**	
Contains			

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: 7/2023 Westminster Pharmaceuticals, LLC