TUSSIN DM- dextromethorphan hbr, doxylamine succinate solution CVS Pharmacy

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

CVS Pharmacy, Inc. Tussin DM Drug Facts

Active ingredients (in each 20 mL)

Dextromethorphan HBr, USP 30 mg

Doxylamine succinate, USP 12.5 mg

Purposes

Cough suppressant

Antihistamine

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat
- controls the impulse to cough to help you sleep

Warnings

Do not use

- to make a child sleepy
- 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

- trouble urinating due to an enlarged prostate gland
- glaucoma
- cough that occurs with too much phlegm (mucus)
- a breathing problem such as emphysema or chronic bronchitis
- persistent or chronic cough such as occurs with smoking, asthma, or emphysema

Ask a doctor or pharmacist before use if you are

taking sedatives or tranquilizers

When using this product

- do not use more than directed
- marked drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

Stop use and ask a doctor if

cough lasts for more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. A persistent cough may be a sign of a serious condition.

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 (1-800-222-1222).

Directions

- measure only with dosing cup provided
- keep dosing cup with product
- mL = milliliter
- do not take more than 4 doses in any 24-hour period
- this adult product is not intended for use in children under 12 years of age

age	dose	
adults and children 12 years and over	20 mL every 6 hours	
children under 12 years	do not use	

Other information

- each 20 mL contains: sodium 11 mg
- store at 20-25°C (68-77°F)

Inactive ingredients

anhydrous citric acid, benzoic acid, benzyl alcohol, carboxymethylcellulose sodium, FD&C blue #1, FD&C red #40, flavor, glycerin, menthol, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose, xanthan gum

Questions or comments?

1-800-719-9260

Principal Display Panel

Compare to the active ingredients in Robitussin® Maximum Strength

Nighttime Cough DM

MAXIMUM STRENGTH

MAXIMUM STRENGTH

Nighttime

 $Tussin\,DM$

DEXTROMETHORPHAN HBr Cough suppressant

DOXYLAMINE SUCCINATE Antihistamine

Adult Cough & Antihistamine

Relieves:

Cough

Itchy throat

Runny nose

Alcohol free

SEE NEW DOSING

Raspberry, Blackberry & Menthol Flavor

For Ages 12 & Over

Dosage cup provided

Actual Bottle Size on Side Panel

4 FL OZ (118 mL)



TUSSIN DM

dextromethorphan hbr, doxylamine succinate solution

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:69842-699 Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9 D2RTI9 KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 20 mL
DOXYLAMINE SUCCINATE (UNII: V9BI9B5YI2) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 20 mL

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)		
BENZOIC ACID (UNII: 8 SKN0 B0 MIM)		
BENZYL ALCOHOL (UNII: LKG8494WBH)		
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679 OBS 311)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
GLYCERIN (UNII: PDC6 A3C0 OX)		
MENTHOL (UNII: L7T10 EIP3A)		
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SORBITOL (UNII: 506T60A25R)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		
XANTHAN GUM (UNII: TTV12P4NEE)		

Product Characteristics			
Color	RED	Score	
Shape		Size	
Flavor	FRUIT	Imprint Code	
Contains			

l	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:69842-699-26	1 in 1 CARTON	06/25/2018	
l	1	118 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	06/25/2018	

Labeler - CVS Pharmacy (062312574)

Revised: 10/2018 CVS Pharmacy