# ADULT TUSSIN DM- dextromethorphan hydrobromide, guaifenesin solution Publix Super Markets Inc

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# Publix Super Markets, Inc. Adult Tussin DM Drug Facts

## Active ingredients (in each 20 mL)

Dextromethorphan HBr, USP 20 mg Guaifenesin, USP 200 mg

## **Purposes**

Cough suppressant Expectorant

#### Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

## **Warnings**

#### Do not use

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

- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis, or emphysema

# 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**

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided
- keep dosing cup with product
- mL = milliliter
- 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 4 hours	
children under 12 years	do not use	

#### Other information

- each 20 mL contains: sodium 14 mg
- store at 68-77°F (20-25°C). Do not refrigerate.

## **Inactive ingredients**

anhydrous citric acid, FD&C red no. 40, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose, xanthan gum

# Package/Label Principal Display Panel

P

**ADULT** 

tussin DM

COUGH SUPPRESSANT (DEXTROMETHORPHAN HBr)

**EXPECTORANT (GUAIFENESIN)** 

Relieves:

- Cough
- Mucus

FOR AGES 12 & OVER

**NON-DROWSY** 

SEE NEW DOSING

Compare to the active ingredients in Robitussin $^{\circledR}$  Cough + Chest Congestion DM 4 FL OZ (118 mL)





COUGH SUPPRESSANT (DEXTROMETHORPHAN HBr) EXPECTORANT (GUAIFENESIN)



Purposes

...Cough suppressant ...Expectorant

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Drug Facts (continued)

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Inactive ingredients anhydrous citric acid, FD&C red no. 40, flavor, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose, xanthan gum

\*This product is not manufactured or distributed by Pfizer, distributor of Robitussin® Cough + Chest Congestion DM.

DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING

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www.StopMedicineAbuse.org



**ADULT** 

# tussinDM

COUGH SUPPRESSANT (DEXTROMETHORPHAN HBr) EXPECTORANT (GUAIFENESIN)



#### Relieves:

- Cough Mucus
- FOR AGES 12 & OVER NON-DROWSY SEE NEW DOSING

\*Compare to the active ingredients in Robitussin® Cough + Chest Congestion DM

4 FL OZ (118 mL)



NDC 56062-607-26

# tussin DM

COUGH SUPPRESSANT (DEXTROMETHORPHAN HBr) EXPECTORANT (GUAIFENESIN)



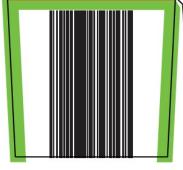
#### Relieves:

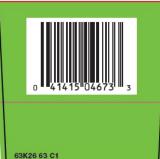
- Cough
- Mucus

FOR AGES 12 & OVER NON-DROWSY SEE NEW DOSING

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4 FL OZ (118 mL)





## **ADULT TUSSIN DM**

dextromethorphan hydrobromide, guaifenesin solution

#### **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:56062-607 **Route of Administration ORAL**

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 20 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)		
SORBITOL (UNII: 506T60A25R)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		
XANTHAN GUM (UNII: TTV12P4NEE)		

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

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56062-607- 26	1 in 1 CARTON	07/21/2022	
1		118 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:56062-607- 34	1 in 1 CARTON	07/28/2022	
2		237 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 Drug	M012	07/21/2022	

Revised: 11/2024 Publix Super Markets Inc