

**TUSSIN DM COUGH AND CHEST CONGESTION- dextromethorphan hbr,  
guaifenesin solution  
NuCare Pharmaceuticals, Inc.**

*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.*

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**Perrigo Tussin DM Drug Facts**

**Active ingredients (in each 10 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	10 mL every 4 hours
children under 12 years	do not use

**Other information**

- Phenylketonurics: Contains Phenylalanine 30 mg per 10 mL
- store at 20-25°C (68-77°F)

**Inactive ingredients**

aspartame, benzoic acid, flavor, glycerin, hydroxyethyl cellulose, menthol, polyethylene glycol, purified water

**Questions or comments?**

1-800-719-9260

**Principal Display Panel**

# NuCare Pharmaceuticals, Inc.

NDC: 68071-4884-4  
Tussin DM 20mg/200mg/10mL

4oz Liquid

See manufacturer's label  
for full list of ingredients.

Product #: R0812004

Tussin DM 20mg/200mg/10mL

Lot: 000000 NDC: 68071-4884-04

MFR NDC: 0113-0578-26 Exp.: 00-00

Tussin DM 20mg/200mg/10mL

Lot: 000000 NDC: 68071-4884-04

MFR NDC: 0113-0578-26 Exp.: 00-00



GTIN 00368071488449

Serial# 00000000002

Exp. Date 00-00

LOT#: 000000

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

Packaged By:  
NuCare Pharmaceuticals, Inc.  
Orange, CA 92867

Distributed by:  
Perrigo Allegan, MI 49010

Take \_\_\_\_\_ teaspoonful(s) every \_\_\_\_\_ hours \_\_\_\_\_ times a day.



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Rev 01/01/19

WARNING: KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED TEMPERATURE 68-77°F.

## TUSSIN DM COUGH AND CHEST CONGESTION

dextromethorphan hbr, guaifenesin solution

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-4884(NDC:0113-0578)
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 10 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 10 mL

### Inactive Ingredients

Ingredient Name	Strength
ASPARTAME (UNII: Z0H242BBR1)	
BENZOIC ACID (UNII: 8SKN0B0MIM)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL (UNII: L7T10EIP3A)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
WATER (UNII: 059QF0KO0R)	

### Product Characteristics

Color		Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-4884-4	118 mL in 1 BOX; Type 0: Not a Combination Product	05/06/2019	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	09/20/1994	

**Labeler** - NuCare Pharmaceuticals,Inc. (010632300)**Establishment**

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals,Inc.		010632300	relabel(68071-4884)

Revised: 2/2021

NuCare Pharmaceuticals,Inc.