# TOPCARE TUSSIN DM- dextromethorphan hydrobromide, guaifenesin solution Topco Associates LLC

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## Topco Associates LLC. 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 20-25°C (68-77°F). 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

#### **Questions or comments?**

1-888-423-0139

## Package/Label Principal Display Panel

TopCare<sub>®</sub> health

COMPARE TO ROBITUSSIN® COUGH + CHEST CONGESTION DM ACTIVE INGREDIENTS

**PEAK COLD** 

SEE NEW DOSING

**NON-DROWSY** 

Tussin DM

Cough & Chest Congestion

COUGH SUPPRESSANT - DEXTROMETHORPHAN HBr

**EXPECTORANT - GUAIFENESIN** 

**RELIEVES:** 

- Cough
- Mucus

#### Adult

For Ages 12 & Over

4 FL OZ (118 mL) RASPBERRY FLAVOR



## **TOPCARE TUSSIN DM**

dextromethorphan hydrobromide, guaifenesin solution

| Product Information     |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:76162-600 |
| 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<br>HYDROBROMIDE | 20 mg<br>in 20 mL  |  |
| GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)                               | GUAIFENESIN                      | 200 mg<br>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<br>Date | Marketing End<br>Date |  |
| 1 | NDC:76162-600-<br>26 | 1 in 1 CARTON   | 05/11/2022              |                       |  |
| 1 |                      | 118 mL in 1 BOTTLE; Type 0: Not a Combination Product |                         |                       |  |
| 2 | NDC:76162-600-<br>34 | 1 in 1 CARTON   | 05/20/2022              |                       |  |
| 2 |                      | 237 mL in 1 BOTTLE; Type 0: Not a Combination Product |                         |                       |  |
|   |                      |   |                         |                       |  |

| Marketing Information |   |                         |                       |  |
|-----------------------|---|-------------------------|-----------------------|--|
| Marketing<br>Category | Application Number or Monograph<br>Citation | Marketing Start<br>Date | Marketing End<br>Date |  |
| OTC Monograph Drug    | M012  | 05/11/2022              |                       |  |
|                       |   |                         |                       |  |

# Labeler - Topco Associates LLC (006935977)

Revised: 11/2024 Topco Associates LLC