# HARRIS TEETER TUSSIN- dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride solution Harris Teeter, LLC

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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# Harris Teeter, LLC Tussin Drug Facts

## Active ingredients (in each 10 mL)

Dextromethorphan HBr, USP 20 mg Guaifenesin, USP 200 mg Phenylephrine HCl, USP 10 mg

#### Purposes

Cough suppressant Expectorant Nasal decongestant

#### Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes
- temporarily relieves these symptoms occurring with a cold:
- nasal congestion
- cough due to minor throat and bronchial irritation

#### 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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema

# When using this product

do not use more than directed

#### Stop use and ask a doctor if

- you get nervous, dizzy, or sleepless
- symptoms do not get better within 7 days or are accompanied by fever
- 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

- each 10 mL contains: sodium 5 mg
- store at 20-25°C (68-77°F).

Do not refrigerate.

#### **Inactive ingredients**

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

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

1-800-719-9260

#### **Principal Display Panel**

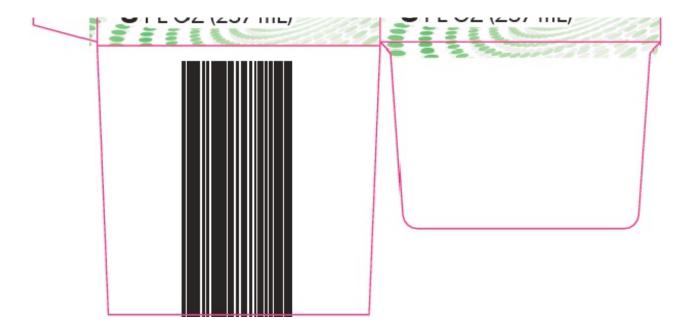
Compare to the active ingredients in Robitussin® Multi-Symptom Cold

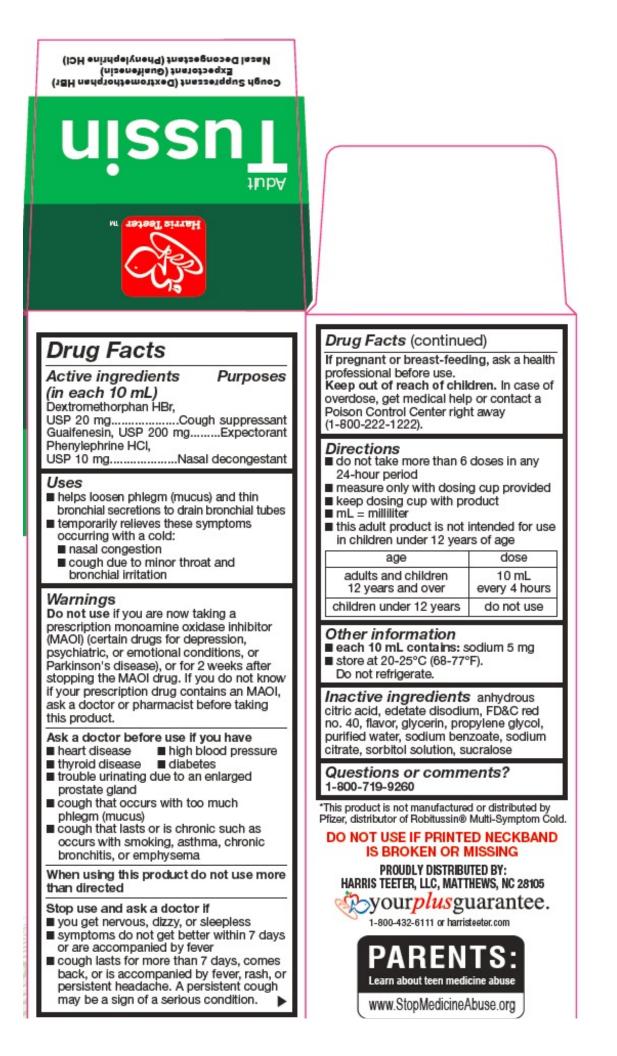
Adult

Tussin

Cough Suppressant (Dextromethorphan HBr) Expectorant (Guaifenesin) Nasal Decongestant (Phenylephrine HCl) MULTI-SYMPTOM COLD Relieves: Cough Nasal Congestion Nasal Congestion Mucus Non-Drowsy CF For Ages 12 & Over 8 FL OZ (237 mL)







| 0 72036 71476 3 |  |
|-----------------|--|
| LOT NO.         |  |
| EXP.            |  |

∙53634 HT C2

| Product Information  |                                 |                    |                                |               |                    |
|--|---------------------------------|--------------------|--------------------------------|---------------|--------------------|
| Product T ype  | HUMAN OTC DRUG                  | Item Code (Source) |                                | NDC:69256-516 |                    |
| Route of Administration  | ORAL                            |                    |                                |               |                    |
| Active Ingredient/Active Mo  | iety                            |                    |                                |               |                    |
| U U  | redient Name                    |                    | Basis of Stre                  | ength         | Strength           |
| DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9 D2RTI9 KYH)<br>(DEXTROMETHORPHAN - UNII:7355X3ROTS) |                                 |                    |                                |               | 20 mg<br>in 10 mL  |
| GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)                             |                                 |                    | GUAIFENESIN                    |               | 200 mg<br>in 10 mL |
| PHENYLEPHRINE HYDRO CHLORII<br>UNII:1WS297W6MV)  | DE (UNII: 04JA59TNSJ) (PHENYLEP | HRINE -            | PHENYLEPHRINE<br>HYDROCHLORIDE |               | 10 mg<br>in 10 mL  |
|  |                                 |                    |                                |               |                    |
| Inactive Ingredients   |                                 |                    |                                |               |                    |
|  | Ingredient Name                 |                    |                                | St            | trength            |
| ANHYDRO US CITRIC ACID (UNII: X  |                                 |                    |                                |               |                    |
| EDETATE DISO DIUM (UNII: 7FLD91  |                                 |                    |                                |               |                    |
| <b>FD&amp;C RED NO. 40</b> (UNII: WZB9127X<br><b>GLYCERIN</b> (UNII: PDC6A3C0OX)           | .UA)                            |                    |                                |               |                    |
| PROPYLENE GLYCOL (UNII: 6DC9   | $\cap 167V3)$                   |                    |                                |               |                    |
| WATER (UNII: 059QF0KO0R)   | Q107 (0)                        |                    |                                |               |                    |
| SODIUM BENZOATE (UNII: OJ245FI   | E5EU)                           |                    |                                |               |                    |
| SODIUM CITRATE (UNII: 1Q73Q2JU   |                                 |                    |                                |               |                    |
| SORBITOL (UNII: 506T60A25R)  |                                 |                    |                                |               |                    |
| SUCRALOSE (UNII: 96K6UQ3ZD4)   |                                 |                    |                                |               |                    |

| Product Characte      | eristic   | s  |                      |                    |  |  |  |
|-----------------------|-----------|--|----------------------|--------------------|--|--|--|
| Color                 |           | RED (clear)                                    | Score                |                    |  |  |  |
| Shape                 |           |  | Size                 |                    |  |  |  |
| Flavor                |           | FRUIT (Menthol)                                | Imprint Code         |                    |  |  |  |
| Contains              |           |  |                      |                    |  |  |  |
|                       |           |  |                      |                    |  |  |  |
|                       |           |  |                      |                    |  |  |  |
| Packaging             |           |  |                      |                    |  |  |  |
| # Item Code           |           | Package Description                            | Marketing Start Date | Marketing End Dat  |  |  |  |
| 1 NDC:69256-516-34    | 1 in 1 C. | ARTON  | 11/21/2015           |                    |  |  |  |
| 1                     | 237 mL    | in 1 BOTTLE; Type 0: Not a Combination Product | t                    |                    |  |  |  |
|                       |           |  |                      |                    |  |  |  |
|                       |           |  |                      |                    |  |  |  |
| Marketing Information |           |  |                      |                    |  |  |  |
| Marketing Categor     | y Ap      | plication Number or Monograph Citation         | Marketing Start Date | Marketing End Date |  |  |  |
| OTC monograph final   | part3     | 41   | 11/21/2015           |                    |  |  |  |
|                       |           |  |                      |                    |  |  |  |

# Labeler - Harris Teeter, LLC (048463103)

Revised: 12/2019

Harris Teeter, LLC