**DIABETIC TUSSIN DM MAXIMUM STRENGTH- dextromethorphan hydrobromide and guaifenesin liquid**
**MEDTECH PRODUCTS 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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**DiabeticTussin DM Max Strength 61787-515**

## Drug Facts

### Active ingredients (in each 10 mL)

Dextromethorphan hydrobromide 20 mg

Guaifenesin 400 mg

### Purposes

Cough Suppressant

Expectorant

### Uses

- temporarily relieves cough caused by the common cold or inhaled irritants.
- helps loosen phlegm (mucus) and thin bronchial secretions to rid bronchial passageways of bothersome mucus.

### 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 two 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

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

### Stop use and ask a doctor if

- cough lasts more than 7 days, comes back, or occurs with fever, rash, or headache that lasts

These could be signs 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.

**Directions**

- take every 4 hours
- do not exceed 6 doses in 24 hours

| | |
|---|---|
| Adults | 10 mL (2 teaspoons) |
| Children under 12 years | ask a doctor |

**Other information**

- store at room temperature 20-25ºC (68-77ºF)
- keep tightly closed

**Inactive ingredients**

Acesulfame K, artificial raspberry flavors, hypromellose, menthol, methylparaben, polyethylene glycol, potassium sorbate, purified water and sucralose. Citric acid may be used to adjust pH.

**Questions or comments?**

Call **1-800-579-8327,** serious side effects associated with use of this product may be reported to this number.

**Package Principal Display Panel**

**SUGAR & ALCOHOL FREE!**

**Specifically Formulated for Diabetics**

**Diabetic Tussin®**

**MAXIMUM STRENGTH**

**COUGH & CHEST CONGESTION**

**DM**

Dextromethorphan HBr (Cough Suppressant)

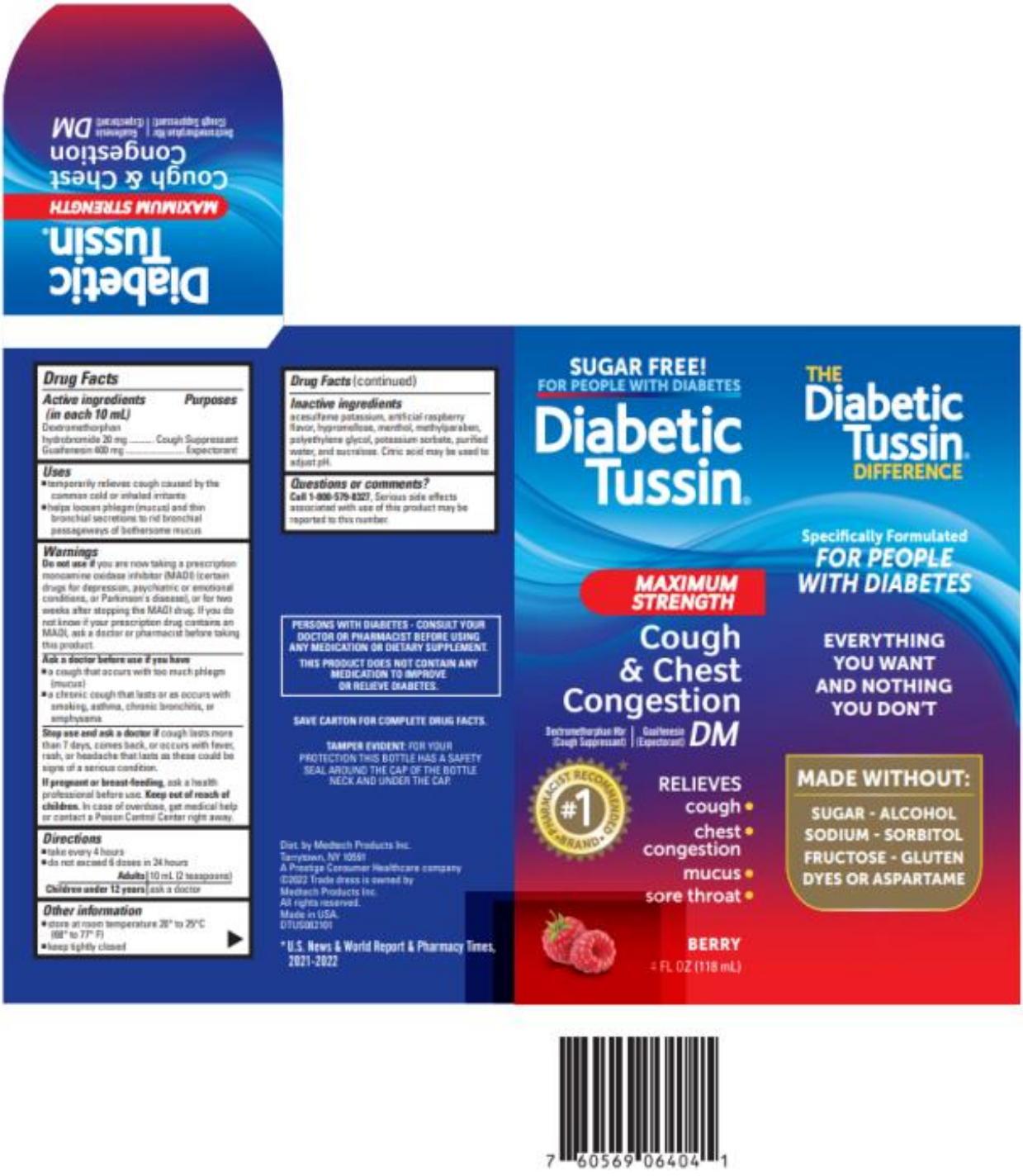
Guaifenesin (Expectorant)

**Relieves:**

- Mucus
- Cough
- Sore throat
- Chest congestion

Improved Berry Flavor

4 FL OZ (118 mL)




## DIABETIC TUSSIN DM MAXIMUM STRENGTH

dextromethorphan hydrobromide and guaifenesin liquid

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:61787-515 |
| **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 | 400 mg in 10 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **ACESULFAME POTASSIUM** (UNII: 23OV73Q5G9) | |
| **HYPROMELLOSE, UNSPECIFIED** (UNII: 3NXW29V3WO) | |
| **MENTHOL, UNSPECIFIED FORM** (UNII: L7T10EIP3A) | |
| **METHYLPARABEN** (UNII: A2I8C7HI9T) | |
| **POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WQ0SDW1A) | |
| **POTASSIUM SORBATE** (UNII: 1VPU26JZZ4) | |
| **RASPBERRY** (UNII: 4N14V5R27W) | |
| **SUCRALOSE** (UNII: 96K6UQ3ZD4) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **ANHYDROUS CITRIC ACID** (UNII: XF417D3PSL) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | | **Score** | |
| **Shape** | | **Size** | |
| **Flavor** | RASPBERRY (artificial raspberry flavor) | **Imprint Code** | |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:61787-515-04 | 1 in 1 BOX | 02/01/2020 | |
| **1** | | 118 mL in 1 BOTTLE; Type 0: Not a Combination Product | | |
| **2** | NDC:61787-515-08 | 1 in 1 BOX | 02/01/2020 | |
| **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 final | part341 | 02/01/2020 | |

## Labeler - MEDTECH PRODUCTS INC (114707784)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Akorn Operating Company LLC (dba Akorn) | | 117696873 | manufacture(61787-515) |

Revised: 3/2022 MEDTECH PRODUCTS INC