**UP AND UP MUCUS RELIEF DM- dextromethorphan hydrobromide, guaifenesin tablet, extended release**
**Target Corporation**

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**Target Corporation Mucus Relief DM Drug Facts**

**Active ingredients (in each extended-release tablet)**

Dextromethorphan HBr 30 mg

Guaifenesin 600 mg

**Purposes**

Cough suppressant

Expectorant

**Uses**

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves:
- cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
- the intensity of coughing
- the impulse to cough to help you get to sleep

**Warnings**

**Do not use**

- for children under 12 years of age
- 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**

- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough accompanied by too much phlegm (mucus)

**When using this product**

- do not use more than directed

**Stop use and ask a doctor if**

- cough lasts more than 7 days, comes back, or occurs with fever, rash, or persistent headache. These could be signs of a serious illness.

**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 crush, chew, or break tablet
- take with a full glass of water
- this product can be administered without regard for timing of meals
- adults and children 12 years and older:

  1 or 2 tablets every 12 hours; not more than 4 tablets in 24 hours
- children under 12 years of age: do not use

**Other information**

- **each tablet contains:** magnesium 15 mg
- do not use if printed foil under cap is broken or missing.
- store between 20-25°C (68-77°F)

**Inactive ingredients**

carbomer homopolymer type B, copovidone, D&C yellow #10 aluminum lake, hypromellose, magnesium hydroxide, magnesium stearate, microcrystalline cellulose, silicon dioxide

**Questions?**

**Call 1-888-547-7400**

**Principal Display Panel**

Compare to active ingredients in Mucinex® DM

mucus relief DM

guaifenesin 600 mg

dextromethorphan hydrobromide 30 mg

extended-release tablets

expectorant and cough suppressant

controls cough

thins and loosens mucus

ACTUAL SIZE

12 HOUR

40 EXTENDED-RELEASE TABLETS

40 EXTENDED-RELEASE TABLETS


## UP AND UP MUCUS RELIEF DM

dextromethorphan hydrobromide, guaifenesin tablet, extended release

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:11673-716 |

| | |
|---|---|
| **Route of Administration** | ORAL |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **DEXTROMETHORPHAN HYDROBROMIDE** (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 30 mg |
| **GUAIFENESIN** (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) | GUAIFENESIN | 600 mg |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL OR ALLYL SUCROSE CROSSLINKED)** (UNII: K6MOM3T5YL) | |
| **COPOVIDONE K25-31** (UNII: D9C330MD8B) | |
| **D&C YELLOW NO. 10** (UNII: 35SW5USQ3G) | |
| **HYPROMELLOSE, UNSPECIFIED** (UNII: 3NXW29V3WO) | |
| **MAGNESIUM HYDROXIDE** (UNII: NBZ3QY004S) | |
| **MAGNESIUM STEARATE** (UNII: 70097M6I30) | |
| **MICROCRYSTALLINE CELLULOSE** (UNII: OP1R32D61U) | |
| **SILICON DIOXIDE** (UNII: ETJ7Z6XBU4) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | YELLOW | **Score** | no score |
| **Shape** | OVAL | **Size** | 16mm |
| **Flavor** | | **Imprint Code** | L219;600 |
| **Contains** | | | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:11673-716-01 | 1 in 1 CARTON | 07/15/2019 | |
| **1** | | 20 in 1 BOTTLE; Type 0: Not a Combination Product | | |
| **2** | NDC:11673-716-49 | 1 in 1 CARTON | 07/15/2019 | |
| **2** | | 40 in 1 BOTTLE; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| ANDA | ANDA207602 | 07/15/2019 | |

## Labeler - Target Corporation (006961700)