

**UP AND UP MUCUS RELIEF- guaifenesin tablet, multilayer, extended release**  
**Target Corporation**

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**Target Corporation Mucus Relief Drug Facts**

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

Guaifenesin 600 mg

**Purpose**

Expectorant

**Uses**

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive

**Warnings**

**Do not use**

- for children under 12 years of age

**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)

**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 overdoses, get medical help or contact a Poison Control Center right away (1-800-222-1222).

**Directions**

- do not crush, chew, or break extended-release tablet

- take with a full glass of water
- this product can be administered without regard for timing of meals
- adults and children 12 years of age and over: 1 or 2 extended-release tablets every 12 hours. Do not exceed 4 extended-release tablets in 24 hours.
- children under 12 years of age: do not use

### **Other information**

- store between 20-25°C (68-77°F)

### **Inactive ingredients**

carbomer homopolymer type B, FD&C blue #1 aluminum lake, hypromellose, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

### **Questions or comments?**

**1-888-547-7400**

### **Package/Label Principal Display Panel**

Compare to active ingredient in Mucinex<sup>®</sup>

Mucus Relief

Guaifenesin Extended-Release Tablets 600 mg

Expectorant

- Relieves chest congestion
- Thins and loosens mucus
- Immediate and extended release

For Day or Night

12 HOURS

up&up<sup>™</sup>

Actual Size

20 Tablets

20 EXTENDED-RELEASE TABLETS



## UP AND UP MUCUS RELIEF

guaifenesin tablet, multilayer, extended release

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11673-421
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	600 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED)</b> (UNII: HHT01Z NK31)	
<b>FD&amp;C BLUE NO. 1 ALUMINUM LAKE</b> (UNII: J9EQA3S2JM)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	

**MAGNESIUM STEARATE** (UNII: 70097M6I30)  
**MICROCRYSTALLINE CELLULOSE** (UNII: OP1R32D61U)  
**SODIUM STARCH GLYCOLATE TYPE A** (UNII: H8AV0SQX4D)

**Product Characteristics**

<b>Color</b>	BLUE	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	16mm
<b>Flavor</b>		<b>Imprint Code</b>	L2X2
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-421-60	1 in 1 CARTON	07/31/2020	02/28/2026
1		20 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11673-421-01	1 in 1 CARTON	09/17/2025	
2		20 in 1 BLISTER PACK; Type 0: Not a Combination Product		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078912	07/31/2020	

**Labeler** - Target Corporation (006961700)