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

Target Corporation Mucus Relief Drug Facts

Active ingredient (in each extended-release tablet)

Guaifenesin 1200 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 overdose, 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 extended-release tablet every 12 hours. Do not exceed 2 extended-release tablets in 24 hours.
- children under 12 years of age: do not use

Other information

store at 20-25°C (68-77°F)

Inactive ingredients

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

Questions or comments?

1-888-547-7400

Package/Label Principal Display Panel

Compare to active ingredient in Maximum Strength Mucinex®

Maximum Strength

Mucus Relief

Guaifenesin Extended-Release Tablets 1,200 mg

Expectorant

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

For Day or Night

12 Hours

up & up™

Actual Size

28 Tablets

28 EXTENDED-RELEASE TABLETS



UP AND UP MUCUS RELIEF

guaifenesin tablet, multilayer, extended release

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82442-503	
Route of Administration	ORAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	1200 mg		

Inactive Ingredients		
Ingredient Name	Strength	
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: HHT01ZNK31)		
COPOVIDONE K25-31 (UNII: D9C330MD8B)		
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		

MAGNESIUM STEARATE (UNII: 70097M6I30)

MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)

Product Characteristics				
Color	BLUE	Score	no score	
Shape	OVAL	Size	22mm	
Flavor		Imprint Code	L4S1	
Contains				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-503- 01	4 in 1 CARTON	09/17/2025	
1		7 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:82442-503- 74	2 in 1 CARTON	09/17/2025	
2		7 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	09/17/2025		

Labeler - Target Corporation (006961700)

Revised: 9/2025 Target Corporation