UP AND UP MUCUS RELIEF- guaifenesin tablet, 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 tablet
- take with a full glass of water
- this product can be administered without regard for the timing of meals
- adults and children 12 years of age and over: 1 tablet every 12 hours. Do not exceed 2 tablets in 24 hours.
- children under 12 years of age: do not use

Other information

- store between 20˚-25˚C (68˚-77˚F)
- do not use if printed foil under cap is broken or missing

Inactive ingredients

colloidal silicon dioxide, copovidone, FD&C blue #1 aluminum lake, hypromellose, magnesium stearate, maltodextrin, microcrystalline cellulose, povidone, sodium starch glycolate Type A, stearic acid

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, 1200 mg

expectorant

relieves chest congestion

thins land loosens mucus

12 HOUR

ACTUAL SIZE

28 TABLETS

28 EXTENDED-RELEASE TABLETS

UP AND UP MUCUS RELIEF

guaifenesin tablet, extended release

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-325
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**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
COPOVIDONE K25-31 (UNII: D9C330MD8B)	

FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	BLUE (Light Blue)	**Score**	no score
Shape	OVAL (Biconvex)	**Size**	22mm
Flavor		**Imprint Code**	Watson;1200
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-325-66	1 in 1 CARTON	07/11/2016	11/30/2019
1		14 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11673-325-30	1 in 1 CARTON	07/11/2016	07/02/2018
2		28 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	ANDA091009	07/11/2016	

Labeler - Target Corporation (006961700)

Revised: 11/2019 Target Corporation