MUCUS RELIEF- mucus relief tablet, extended release
Allegiant Health

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459 - Mucus Relief

Active ingredient(s)

Guaifenesin 600 mg

Purpose

Expectorant

Use(s)

- 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

- 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 breastfeeding,

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 of age and over: 1 or 2 tablets every 12 hours. Do not exceed 4 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 imprinted safety seal under cap is broken or missing

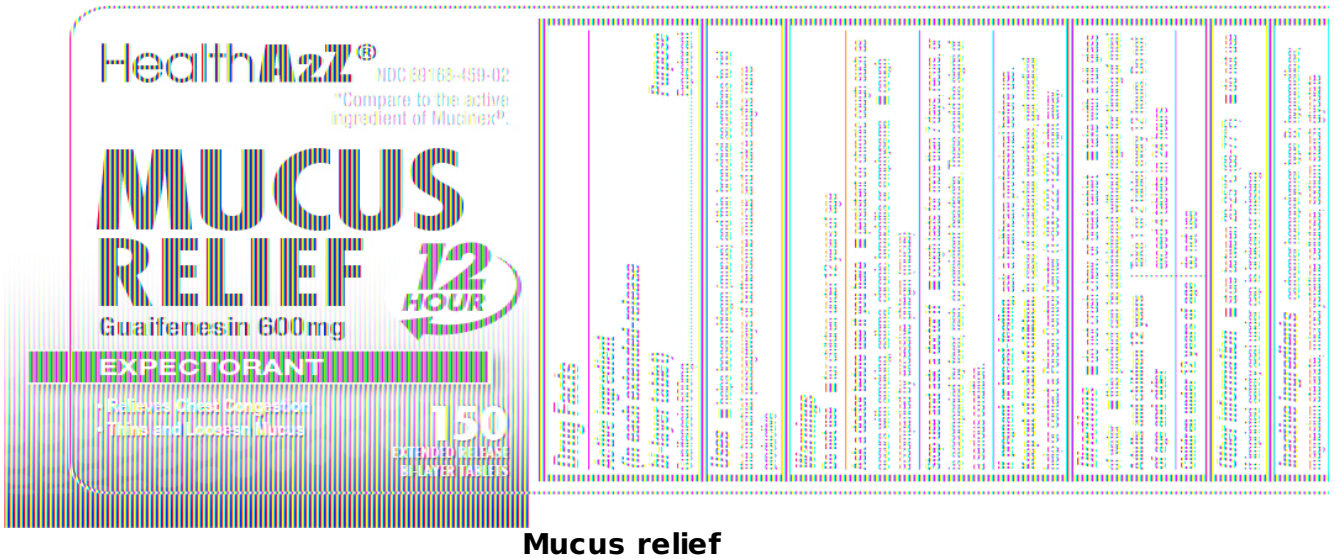
Inactive ingredients

carbomer homopolymer type B; hypromellose, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

Questions/Comments

Call 1-888-952-0050 Monday through Friday 9AM – 5PM EST.

Principal Display Panel



Mucus relief

MUCUS RELIEF			
mucus relief tablet, extended release			
Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69168-459
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
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)	
HYPROMELLOSE 2910 (5 MPA.S) (UNII: R75537T0T4)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

Product Characteristics

Color	white	**Score**	no score
Shape	OVAL	**Size**	16mm
Flavor		**Imprint Code**	G;600
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69168-459-75	75 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2024	
2	NDC:69168-459-02	150 in 1 BOTTLE; Type 0: Not a Combination Product	04/24/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA213420	04/24/2024	

Labeler - Allegiant Health (079501930)

Revised: 4/2024 Allegiant Health