MUCUS RELIEF- guaifenesin tablet, extended release
Advanced Rx LLC

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

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 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 20º to 25ºC (68º to 77ºF)

■ DO NOT USE IF PRINTED SEAL UNDER CAP IS MISSING OR DAMAGED

Inactive ingredients

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

Questions?

Call toll-free **1-800-630-8895**

Distributed by:

ADVANCED RX LLC

1942 NE 163rd St North Miami Beach,

FL 33162 U.S.A

PRINCIPAL DISPLAY PANEL

NDC 80513-403-20

*Compare to the active ingredient in Mucinex ®

Mucus Relief

Guaifenesin 600 mg

Expectorant

EXTENDED-RELEASE TABLETS BI-LAYER TABLETS

200 Tablets

*This product is not manufactured or distributed by the owner of the registered trademark Mucinex ®Extended-Release 600 mg Tablets

MUCUS RELIEF

guaifenesin tablet, extended release

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:80513-403
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 CROSSLINKED) (UNII: HHT01ZNK31)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
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:80513-403-20	200 in 1 BOTTLE; Type 0: Not a Combination Product	08/29/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA213420	05/08/2020	

Labeler - Advanced Rx LLC (042795108)

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