# MUCUS RELIEF- guaifenesin tablet, extended release Advanced Rx LLC

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# **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 20<sup>o</sup> to 25<sup>o</sup>C (68<sup>o</sup> to 77<sup>o</sup>F)
- 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 <sup>®</sup>Extended-Release 600 mg Tablets

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Product Information								
Product Type	HUMAN OTC DRUG Item Code (Source) NDC:							
Route of Administration	ORAL							
Active Ingredient/Activ	ve Moiety							
GUAIFENESIN (UNII: 495W7451			7451VQ)	GUAIFENESIN	Strength			
CARBOMER HOMOPOLYMER		ngredient Nam LYL PENTAERYTH		INKED) (UNII:		Strengt		
HHI012NK31) HYPROMELLOSE, UNSPECIFII	<b>D</b> (IINII) 3N							
SODIUM STARCH GLYCOLATE								
SODIUM STARCH GLYCOLATE MAGNESIUM STEARATE (UNII:	70097M6I30	)						
MAGNESIUM STEARATE (UNII:								
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MAGNESIUM STEARATE (UNII: MICROCRYSTALLINE CELLULO Product Characteristic	DSE (UNII: C			n	o score			
MAGNESIUM STEARATE (UNII: MICROCRYSTALLINE CELLULO Product Characteristic Color	<b>DSE</b> (UNII: C : <b>S</b>	0P1R32D61U)			o score 6mm			
MAGNESIUM STEARATE (UNII: MICROCRYSTALLINE CELLUL	DSE (UNII: C S white	P1R32D61U) Score	de	1				

Packaging								
#	Item Code	Package Description	Marketing Start Date	Marketing End Date				
	NDC:80513-403- 20	200 in 1 BOTTLE; Type 0: Not a Combination Product	08/29/2024					
Marketing Information								
	Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date				
AN	DA	ANDA213420	05/08/2020					

Labeler - Advanced Rx LLC (042795108)

Revised: 8/2024

Advanced Rx LLC