MUCUS RELIEF- guaifenesin 200 mg tablet
Richmond Pharmaceuticals, Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Guaifenesin Caplets, 200 mg

Active ingredient (in each tablet)

Guaifenesin 200 mg

Purpose

Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive to rid the bronchial passageway of bothersome mucus and make cough more productive

Warnings

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 excessive phlegm (mucus)

KEEP OUT OF REACH OF CHILDREN

In case of overdose, get medical help or contact a Poison Control Center right away.

IN CASE OF PREGNANCY OR BREAST FEEDING SECTION

ask a health professional before use

stop use and ask a doctor

if cough lasts more than 7 days, comes back, or is accompanied by fever, rash or persistent headache.These could be signs of serious illness

Directions:

- **adults and children 12 years of age and over:** take 1 to 2 tablets every 4 hours with a full glass of water while symptoms persist. Do not exceed 6 doses in 24 hours.
- **children under 12 years:** do not use

OTHER INFORMATION

- store at 15°-30°C (59°-86°F)
- TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Inactive ingredients

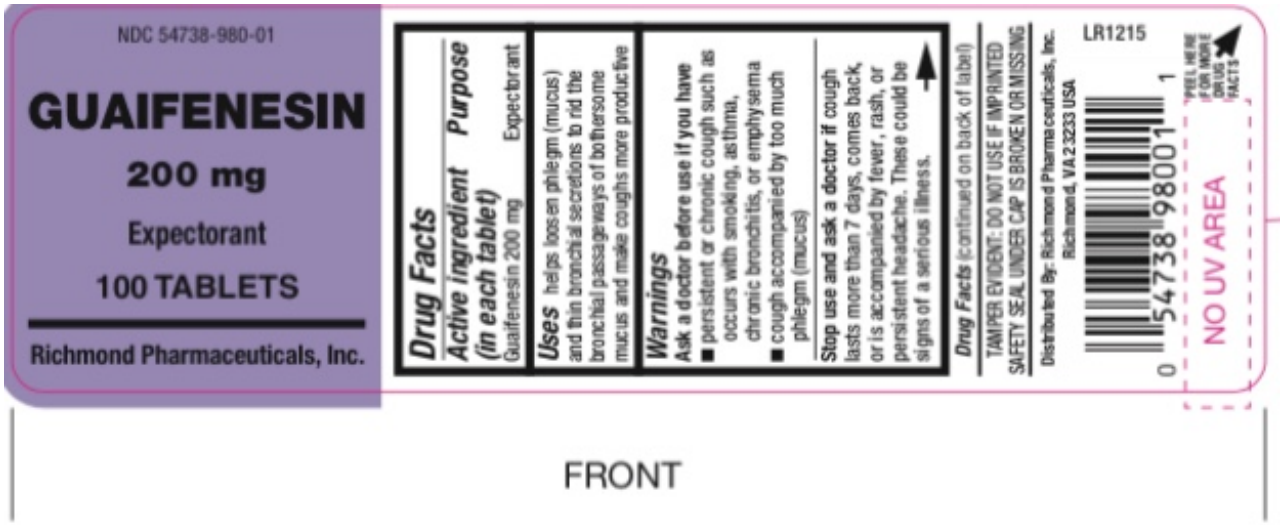
colloidal silicon dioxide, FD&C RED#40(Al-lake), magnesium stearate, maltodextrin, microcrystalline

cellulose, stearic acid, sodium starch glycolate

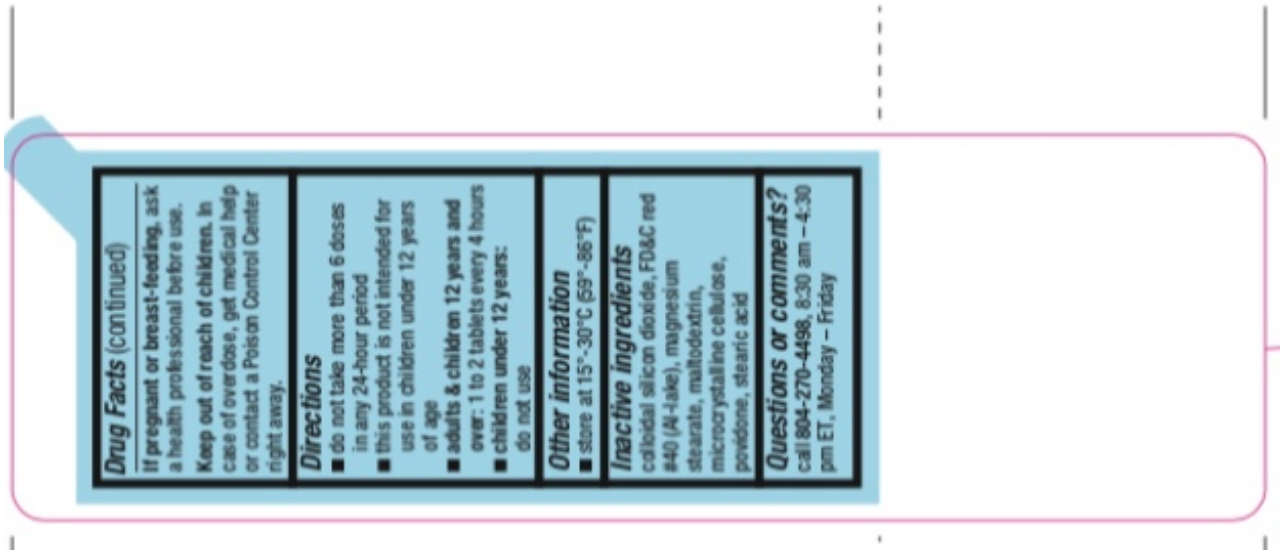
Questions or Comments

call 804-270-4498, 8.30 am-4.30 pm ET, Monday - Friday
TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

Guaifenesin 200 mg...Expectorant
NDC- 54738-980-01... 100 CAPLETS



FRONT



MUCUS RELIEF

guaifenesin 200 mg tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54738-980
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg

Inactive Ingredients	
Ingredient Name	**Strength**
FD&C RED NO. 40 (UNII: WZB9127XOA)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POVIDONE (UNII: FZ989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics			
Color	red	**Score**	2 pieces
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	AP;151
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:54738-980-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2016	

Marketing Information

Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC monograph final	part341	03/01/2016	

Labeler - Richmond Pharmaceuticals, Inc. (043569607)

Registrant - Advance Pharmaceutical Inc. (078301063)

Establishment

Name	**Address**	**ID/FEI**	**Business Operations**
Advance Pharmaceutical Inc.		078301063	manufacture(54738-980)