

MUCUS RELIEF - guaifenesin tablet, extended release
McKesson Corporation dba SKY Packaging

SKY Mucus Relief

ACTIVE INGREDIENT(in each extended-release tablet)

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

WARNING

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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 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

- tamper evident: do not use if printed safety seal is broken or missing
- store between 20 to 25°C (68 to 77°F)

INACTIVE INGREDIENTS

carbomer homopolymer, hypromellose, microcrystalline cellulose, povidone

QUESTIONS?

1-888-759-4633

Hours: 8am - 4pm, EST, MON - FRI.

You may also report side effects to this phone number.

PRINCIPAL DISPLAY PANEL

SKY

NDC 63739-067-02

MUCUS RELIEF

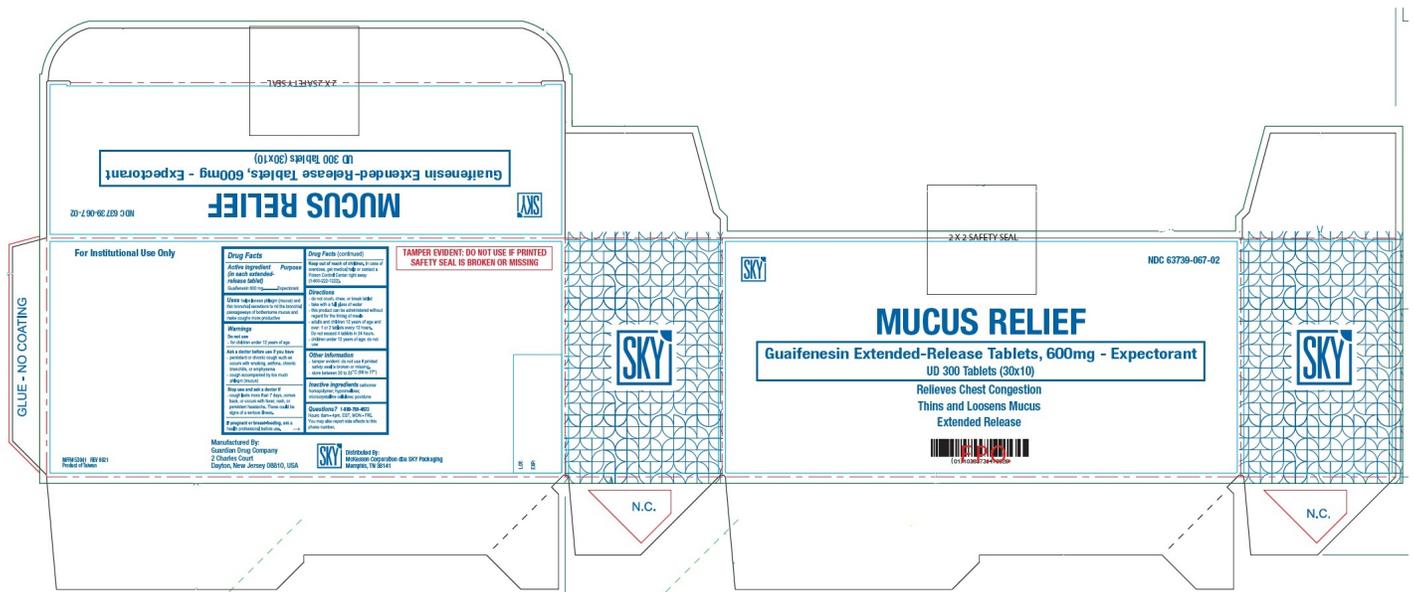
Guaifenesin Extended-Release Tablets, 600 mg - Expectorant

UD 300 Tablets (30x10)

Relieves Chest Congestion

Thins and Loosens Mucus

Extended Release



MUCUS RELIEF

guaifenesin tablet, extended release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63739-067
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, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics

Color	WHITE	Score	no score
Shape	CAPSULE	Size	22mm
Flavor		Imprint Code	G233
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63739-067-02	30 in 1 BOX	10/20/2021	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA209215	10/20/2021	

Labeler - McKesson Corporation dba SKY Packaging (140529962)

Establishment

Name	Address	ID/FEI	Business Operations
Guardian Drug Company		119210276	MANUFACTURE(63739-067)

Revised: 11/2022

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