

SUNMARK MUCUS RELIEF - guaifenesin tablet, extended release
Strategic Sourcing Services LLC

Sunmark mucus relief
Guaifenesin Extended-Release Tablets 600 mg

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 carton is open or if printed seal on blister is broken or missing
- store between 20 to 25°C (68 to 77°F)

INACTIVE INGREDIENTS

carbomer homopolymer, hypromellose, microcrystalline cellulose, povidone

QUESTIONS?

1-609-860-2600

Hours: 8am - 4pm, EST

You may also report side effects to this phone number.

PRINCIPAL DISPLAY PANEL

sunmark

COMPARE TO THE ACTIVE INGREDIENT IN MUCINEX® EXTENDED RELEASE 600 MG TABLETS

NDC 70677-0055-1

12 hour

mucus relief

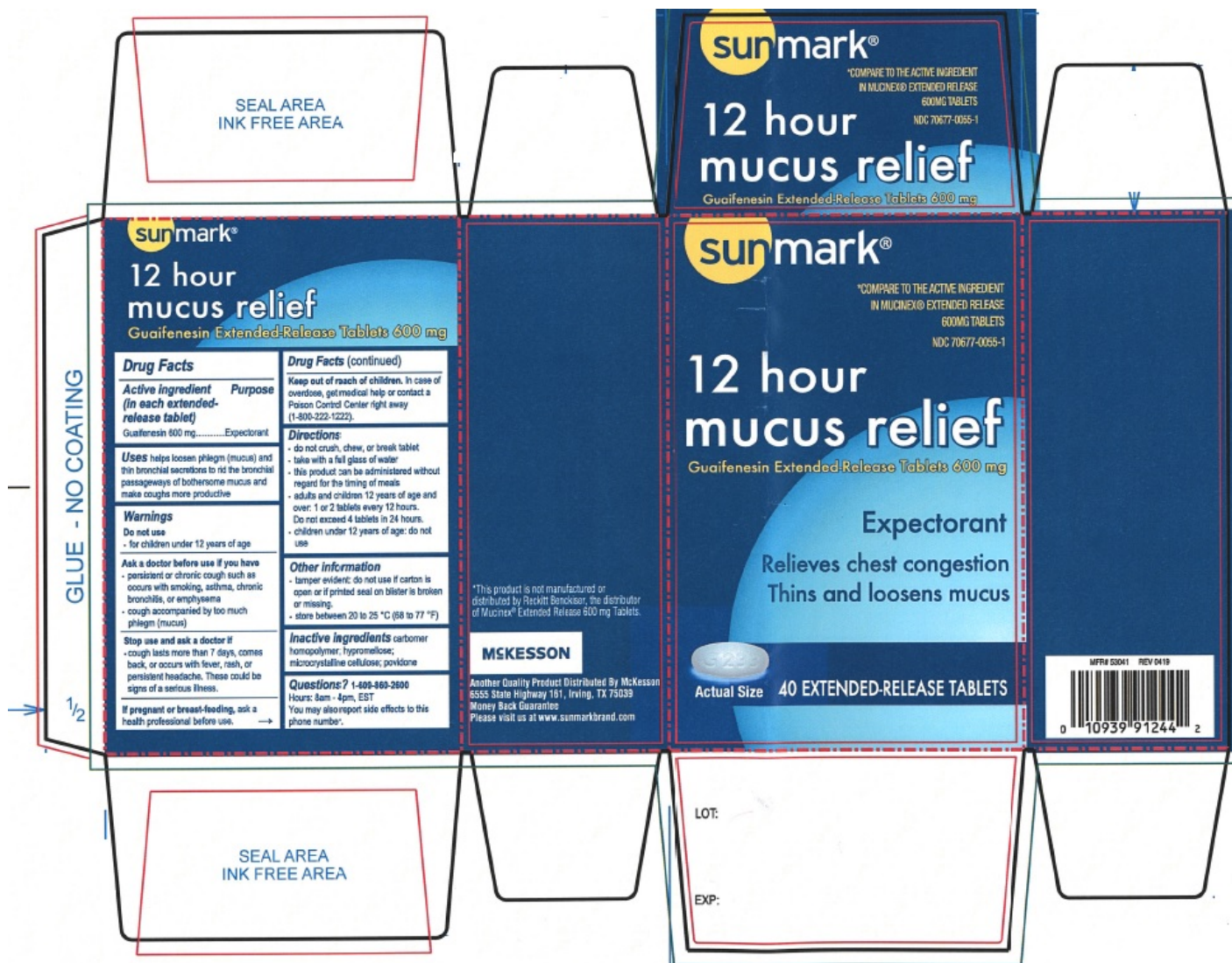
Guaifenesin Extended-Release Tablets 600 mg

Expectorant

Relieves chest congestion

Thins and Loosens Mucus

40 Extended-Release Tablets



SUNMARK MUCUS RELIEF

guaifenesin tablet, extended release

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-0055
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:70677-0055-1	4 in 1 CARTON	09/26/2019	
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	07/22/2018	

Labeler - Strategic Sourcing Services LLC (116956644)

Revised: 1/2022

Strategic Sourcing Services LLC