

GUAIFENESIN 200MG- guaifenesin tablet
HEALTHLIFE OF USA LLC

Guaifenesin 200mg

Active ingredient (in each tablet)

Guaifenesin 200mg

Purpose

Expectorant

Guaifenesin 200 mg Tablets

Uses

helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageway of bothersome mucus and make coughs 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 too much phlegm (mucus)

Stop use and ask a doctor if

cough last more than 7 days, come back or is accompanied by fever, rash, or persistent headache. There could be signs of a serious illness.

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

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.

Directions:

- Do not take more than 6 doses in any 24-hour period.
- This product is not intended for use in children under 12 years of age
- Adults & children 12 years and over: 1 to 2 tablets every 4 hours

- Children under 12 years: do not use

Other Information:

- Store at 15°C-30°C (59°F-86°F)

Inactive Ingredients:

Colloidal silicon dioxide, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

Manufactured & Distributed by:

Health Pharma

Rahway, NJ 07065

www.healthpharma.us

Questions or comments?

Call toll free 1-844-832-1138 Monday through Friday 9AM - 5PM EST

PRINCIPAL DISPLAY PANEL

Guaifenesin 200mg Tablets

NDC 69517-148-03

Count: 300



Drug Facts	Purpose Expectorant
Active ingredient (in each tablet) Guaifenesin 200 mg	
Uses ■ helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive to rid the bronchial passageway of bothersome mucus	
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) When using this product ■ do not exceed recommended dosage ■ do not use for more than 7 days Stop use and ask a doctor if cough lasts for more than 7 days, recurs, or is accompanied by fever, rash, or persistent headache. These could be signs of a serious condition. 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.	
Directions ■ adults and children 12 years of age and over: take 1 tablet 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 ■ each tablet contains: sodium 1.24 mg VERY LOW SODIUM ■ store at 25°C (77°F); excursions between 15°-30°C (59°-86°F) ■ keep in a dry place and do not expose to heat ■ read all product information before using ■ TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING	
Inactive ingredients colloidal silicon dioxide, magnesium stearate, microcrystalline cellulose, sodium starch glycolate	
Questions or comments? Call toll free 1-844-832-1138 Monday through Friday 9AM - 5PM EST or www.healthlifeofusa.com	

Distributed by:
HealthLife® of USA LLC
Rahway, NJ 07065
www.healthlifeofusa.com

MADE IN THE USA

Lot No.:
Exp. Date:

3 69517 14803 3

GUAIFENESIN 200MG guaifenesin tablet
Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69517-148
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
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	G2
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69517-148-03	300 in 1 BOTTLE; Type 0: Not a Combination Product	01/28/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/28/2022	

Labeler - HEALTHLIFE OF USA LLC (079656178)

Establishment

Name	Address	ID/FEI	Business Operations
Health Pharma USA LLC		080804485	manufacture(69517-148)

Revised: 10/2023

HEALTHLIFE OF USA LLC