CURIST MUCUS RELIEF MAXIMUM STRENGTH- guaifenesin tablet, extended release
Little Pharma, Inc.

Curist Mucus Relief Max Str (Guaifenesin 1200 mg) ER

Drug Facts

Active ingredient (in each extended-release tablet)

Guaifenesin 1200 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 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 tablet every 12 hours. Do not exceed 2 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°-25°C (68°-77°F)

Inactive ingredients carbomer homopolymer type B, hypromellose, magnesium

stearate, microcrystalline cellulose, sodium starch glycolate

Questions?

Contact **1-844-243-1241** or

email hi@curistrelief.com

Distributed by:
Little Pharma, Inc.

New York, NY 10023

Made in India

curist

Mucus Relief Maximum Strength

Guaifenesin Extended-Release Tablets 1200 mg

Expectorant

12 Hour

Relieves Chest Congestion

Thins and Loosens Mucus

Immediate and Extended Release

150 Extended-Release Tablets

CURIST MUCUS RELIEF MAXIMUM STRENGTH

guaifenesin tablet, extended release

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72559-013(NDC:62207-840)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	1200 mg

Inactive Ingredients

Ingredient Name	Strength
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
CARBOMER HOMOPOLYMER TYPE B (ALLYL PENTAERYTHRITOL OR ALLYL SUCROSE CROSSLINKED) (UNII: K6MOM3T5YL)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

Product Characteristics

Color	white	**Score**	no score
Shape	OVAL	**Size**	22mm
Flavor		**Imprint Code**	G;1200
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72559-013-11	6 in 1 CARTON	07/23/2021	
1		14 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:72559-013-28	150 in 1 BOTTLE; Type 0: Not a Combination Product	04/06/2022	
3	NDC:72559-013-41	2 in 1 PACKAGE, COMBINATION	07/23/2021	
3		150 in 1 BOTTLE; Type 1: Convenience Kit of Co-Package		
4	NDC:72559-013-64	84 in 1 BOTTLE; Type 0: Not a Combination Product	11/15/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA213420	07/23/2021	

Labeler - Little Pharma, Inc. (074328189)

Revised: 3/2026 Little Pharma, Inc.