MAX TUSSIN MUCUS AND CHEST CONGESTION SUGAR FREEguaifenesin liquid Akron Pharma 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.

Drug Facts

Active ingredients (in each 10 mL)

Guaifenesin 200 mg

Purposes

Expectorant

Uses

 helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive

Warnings

Ask a doctor before use if you have

- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema

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 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 (1-800-222-1222).

Directions

- do not take more than 6 doses in any 24-hour period
- mL= milliliter
- this adult product is not intended for use in children under 12 years of age

age	dose
adults and children 12 years and over	10 - 20 mL every 4 hours
children under 12 years	do not use

Other information

- Phenylketonurics: contains Phenylalanine 30 mg per 10 mL
- store between 20-25°C(68-77°F). Do not refrigerate

Inactive ingredients

anhydrous citric acid, flavor cherry, glycerin, menthol, purified water, saccharin sodium, sodium benzoate, sorbitol solution, sucralose.

Questions or comments?

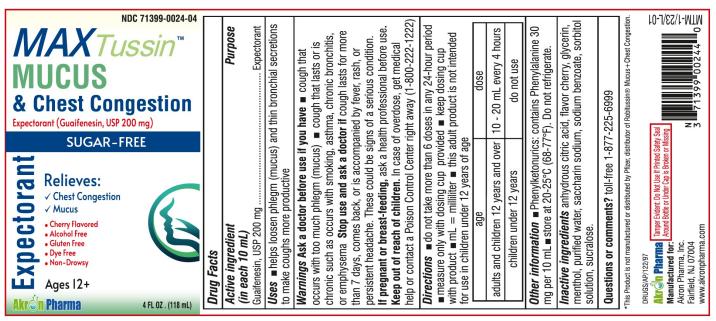
toll-free 1-877-225-6999

Manufactured for:

Akron Pharma, Inc.

Fairfield, NJ 07004

www.akronpharma.com





MUCUS & Chest Congestion

Expectorant (Guaifenesin, USP 200 mg)

SUGAR-FREE

✓ Chest Congestion √ Mucus Alcohol Free Gluten Free Dye Free Non-Drowsy

Relieves:

- Cherry Flavored





8 FL 0Z . (236 mL)

Drug Facts

Active ingredient (in each 10 mL)

Guaifenesin, USP 200 mg Expectorant

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Directions ■ do not take more than 6 doses in any 24-hour period ■ measure only with dosing cup provided ■ keep dosing cup with product

Drug Facts (continued)

Purpose

■mL = milliliter ■this adult product is not intended for use in children under 12 years of age

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Questions or comments? toll-free 1-877-225-6999

*This Product is not manufactured or distributed by Pfizer, distributor of Robitussin®

Tamper Evident: Do Not Use If Printed Safety Seal Around Bottle or Under Cap is Broken or Missing

DRUGS/AP/122/97

Akr@n Pharma

Purpose

Manufactured for: Akron Pharma, Inc. Fairfield, NJ 07004 www.akronpharma.com



NDC 71399-0024-06

16 FL OZ . (473 mL)

MAXTussin™

MUCUS & Chest Congestion

Expectorant (Guaifenesin, USP 200 mg)

SUGAR-FREE



Drug Facts Active ingredient

(in each 10 mL)

Guaifenesin, USP 200 mg Expectorant

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Drug Facts (continued)

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Questions or comments?toll-free 1-877-225-6999

*This Product is not manufactured or distributed by Pfizer, distributor of Robitussin® Mucus + Chest Congestion.

Tamper Evident: Do Not Use If foil seal over bottle opening is torn broken or missing.

DRUGS/AP/122/97

Akran Pharma

Manufactured for: Akron Pharma, Inc. Fairfield, NJ 07004 www.akronpharma.com



MAX TUSSIN MUCUS AND CHEST CONGESTION SUGAR FREE

guaifenesin liquid

Akren Pharma

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71399-0024
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 10 mL

Inactive Ingredients			
Ingredient Name	Strength		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)			
GLYCERIN (UNII: PDC6A3C0OX)			
MENTHOL (UNII: L7T10EIP3A)			
WATER (UNII: 059QF0KO0R)			
SACCHARIN SODIUM (UNII: SB8ZUX40TY)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
SORBITOL SOLUTION (UNII: 8KW3E207O2)			
SUCRALOSE (UNII: 96K6UQ3ZD4)			

Product Characteristics			
Color		Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399- 0024-4	1 in 1 PACKAGE	05/15/2023	
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:71399- 0024-8	1 in 1 PACKAGE	05/15/2023	
2		236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:71399- 0024-6	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/15/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part341	05/15/2023	

Labeler - Akron Pharma Inc. (067878881)

Revised: 5/2023 Akron Pharma Inc.