MAX TUSSIN MUCUS AND CHEST CONGESTION- guaifenesin liquid Akron Pharma Inc.

MAX TUSSIN MUCUS & Chest Congestion Expectorant

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
- measure only with dosing cup provided
- 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 at 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, sucrose.

Questions or comments?

toll-free 1-877-225-6999

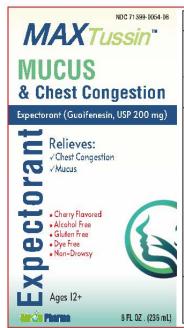
Manufactured for:

Akron Pharma, Inc.

Fairfield, NJ 07004

www.akronpharma.com

150 mm x 76 mm



| Drug Facts | |
|-------------------------|-------------|
| Active ingredient | Purpose |
| (in each 10 mL) | |
| Guaifenesin, USP 200 mg | 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 for 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 ■ measure only with dosing cup provided ■ keep dosing cup with product

Drug Facts (continued)

■ 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 at 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, sucrose

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

*This Product is not manufactured or distributed by Pitzer, distributor of Robitussin® Mucus + Chest Congestion. Tamper Evident: Do Not Use If Printed Safety Seal Around Bottle or Under Cap is Broken or Missing





MAX TUSSIN MUCUS AND CHEST CONGESTION

quaifenesin liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:71399-0054

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

SUCRALOSE (UNII: 96K6UQ3ZD4)
SUCROSE (UNII: C151H8M554)

| 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) | |
| | |

| Product Characteristics | | | |
|-------------------------|--------|--------------|--|
| Color | | Score | |
| Shape | | Size | |
| Flavor | CHERRY | Imprint Code | |
| Contains | | | |

| Packaging | | | | |
|------------|------|---|-------------------------|-----------------------|
| # Item (| Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 NDC:7139 | 99- | 1 in 1 PACKAGE | 11/03/2023 | |
| 1 | | 236 mL in 1 BOTTLE; Type 0: Not a Combination Product | | |

| Marketing Information | | | |
|-----------------------|---|-------------------------|-----------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| OTC Monograph Drug | M012 | 11/03/2023 | |
| | | | |

Labeler - Akron Pharma Inc. (067878881)

Registrant - HI-TECH PHARMACEUTICALS PRIVATE LIMITED (862584260)

| Establishment | | | | |
|---|---------|-----------|----------------------------|--|
| Name | Address | ID/FEI | Business Operations | |
| HI-TECH PHARMACEUTICALS PRIVATE LIMITED | | 862584260 | manufacture(71399-0054) | |

Revised: 11/2023 Akron Pharma Inc.