

**MAX TUSSIN COUGH AND CHEST CONGESTION DM- dextromethorphan hbr,
guaifenesin liquid
Akron Pharma Inc.**

**Max Tussin Cough and Chest Congestion DM
Guaifenesin 200mg and Dextromethorphan Hbr 20mg per 10ml
Alcohol-Free
Cherry Flavor**

Active ingredients (in each 10 mL)

Dextromethorphan HBr 20 mg

Guaifenesin 200 mg

Purposes

Cough suppressant

Expectorant

Uses

- temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

Warnings

Do not use

if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

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

If pregnant or breast-feeding,

ask a health professional before use.

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.

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
- 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 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, sodium citrate, sucralose, sucrose.

Questions or comments?

toll-free 1-877-225-6999

Manufactured for:

Akron Pharma, Inc.

Fairfield, NJ 07004

www.akronpharma.com

MAXTussin™ COUGH & CHEST CONGESTION DM

NDC 71399-0025-4

Cough Suppressant (Dextromethorphan HBr)
Expectorant (Guaifenesin)

Congestion

- ✓ Controls Coughs
- ✓ Loosens & Relieves Chest Congestion

- Cherry Flavored
- Alcohol Free
- Gluten Free
- Dye Free
- Non-Drowsy

Ages 12+



Akron Pharma

4 FL. OZ. (118 mL)

Drug Facts

Active ingredients (in each 10 mL)
Dextromethorphan HBr, USP 20 mg Cough suppressant
Guaifenesin, USP 200 mg Expectorant

Uses ■ temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold ■ helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes

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Other information ■ Each 10mL contains: Sodium 10.23mg ■ store at 20°-25°C (68°-77°F). Do not refrigerate.

Inactive ingredients anhydrous citric acid, flavor cherry, glycerin, menthol, propylene glycol, purified water, sodium benzoate, sodium citrate, sucralose, sucrose

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

*This Product is not manufactured or distributed by Pfizer distributor of Robbussin® Cough+ Chest Congestion DM
Tamper Evident: Do Not Use If Printed Safety Seal Around Bottle or Under Cap is Broken or Missing

LOT No.:
EXPDate:
Un Warnish Area

REV 07/25
N 71399-0025-4 9
www.akronpharma.com

MAXTussin™
COUGH
& CHEST CONGESTION DM

Cough Suppressant (Dextromethorphan HBr)
Expectorant (Guaifenesin)

NDC 71399-0025-4

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NDC 71399-0025-4

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REV 07/25
N 71399-0025-4 9
www.akronpharma.com

Size : 50x130x50mm Lock Bottom

NDC 71399-0025-8

MAXTussin™

COUGH & CHEST CONGESTION DM

Cough Suppressant (Dextromethorphan HBr)
Expectorant (Guaifenesin)

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Ages 12+



8 FL OZ. (236 mL)

Drug Facts

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Drug Facts (Continued)

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 Akron Pharma, Inc.
 Fairfield, NJ 07004
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Un Warnish Area

REV.07/25



Size : 58x163x58mm Lock Bottom

NDC 71399-0025-8

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

Rev.07/25



NDC 71399-0025-6

MAXTussin™

COUGH & CHEST CONGESTION DM

Cough Suppressant (Dextromethorphan HBr)
Expectorant (Guaifenesin)

Congestion

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Ages 12+



16 FL OZ. (473 mL)

Drug Facts

Active ingredients (in each 10 mL)

Dextromethorphan HBr, USP 20 mg .. Cough suppressant
Guaifenesin, USP 200 mg Expectorant

Purposes

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Drug Facts (Continued)

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

Un Warnish Area



Manufactured for:
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Fairfield, NJ 07004
www.akronpharma.com



MAX TUSSIN COUGH AND CHEST CONGESTION DM

dextromethorphan hbr, guaifenesin liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71399-0025
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII: 7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 10 mL
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)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SACCHARIN SODIUM (UNII: SB8ZUX40TY)	

SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-0025-4	1 in 1 PACKAGE	05/15/2023	
1		118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:71399-0025-8	1 in 1 PACKAGE	05/15/2023	
2		236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:71399-0025-6	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 Drug	M012	05/15/2023	

Labeler - Akron Pharma Inc. (067878881)

Revised: 10/2025

Akron Pharma Inc.