

MUCUS RELIEF DM MAX- dextromethorphan hbr, guaifenesin solution
Family Dollar Services Inc

Family Wellness 44-031A

Active ingredients (in each 20 mL)

Dextromethorphan HBr 20 mg
Guaifenesin 400 mg

Purpose

Cough suppressant
Expectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves:
 - cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
 - the intensity of coughing
 - the impulse to cough to help you get to sleep

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

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

Stop use and ask a doctor if

cough persists more than 1 week, tends to recur, or is accompanied by a 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

- **do not take more than directed**
- do not take more than 6 doses in any 24-hour period
- mL = milliliter
- only use the dose cup provided
- dose as follows or as directed by a doctor
- adults and children 12 years and over: 20 mL in dosing cup provided every 4 hours
- children under 12 years: do not use

Other information

- **each 20 mL contains:** sodium 6 mg
- use by expiration date on package
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

Inactive ingredients

anhydrous citric acid, FD&C red #40, flavors, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sorbitol solution, sucralose, xanthan gum

Questions or comments?

1-800-426-9391

Principal display panel

FAMILY
Wellness™

***COMPARE TO THE
ACTIVE INGREDIENTS IN
MAXIMUM STRENGTH
MUCINEX® FAST-MAX®
DM MAX**

MAXIMUM STRENGTH
**MUCUS RELIEF
DM MAX**

**Dextromethorphan HBr
Guaifenesin**

**Cough Suppressant
Expectorant**

- **Controls Cough**

- **Relieves Chest Congestion**
- **Thins & Loosens Mucus**

**AGES
12 YEARS
& OVER**

**6 FL OZ
(177 mL)**

NDC 55319-731-45

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

**PARENTS:
Learn about teen medicine abuse
www.StopMedicineAbuse.org**

*This product is not manufactured or distributed by RB Health (US) LLC, owner of the registered trademark Maximum Strength Mucinex® FAST-MAX® DM MAX.
50844 ORG042403145

**DISTRIBUTED BY:
MIDWOOD BRANDS LLC,
500 VOLVO PKWY
CHESAPEAKE, VA 23320 USA**

NOT 100% SATISFIED?
Return within 30 days to the store of purchase for a refund (with receipt) or exchange.

FAMILY Wellness™ *COMPARE TO THE ACTIVE INGREDIENTS IN MAXIMUM STRENGTH MUCINEX® FAST-MAX® DM MAX

MAXIMUM STRENGTH
MUCUS RELIEF DM MAX

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F-031A-45 ORG NDC 55319-731-45

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PEEL BACK TAB TO READ COMPLETE DRUG FACTS AND INFORMATION

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

Drug Facts

Active ingredients (in each 20 mL)	Purpose
Dextromethorphan HBr 20 mg.....	Cough suppressant
Guaifenesin 400 mg	Expectorant

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SKU 999559

0 32251 92385 2

No print/No varnish
Lot & Exp date

Drug Facts (continued)

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B-031A-45
ORG

Family Wellness 44-031A

MUCUS RELIEF DM MAX

dextromethorphan hbr, guaifenesin solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55319-731
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	red	Score	
Shape		Size	
Flavor	BERRY	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55319-731-45	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/05/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/05/2024	

Labeler - Family Dollar Services Inc (024472631)

Establishment

Name	Address	ID/FEI	Business Operations
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LNK International, Inc.

967626305

manufacture(55319-731) , pack(55319-731)

Revised: 1/2026

Family Dollar Services Inc