

**DM MAX CONGESTION AND COUGH- dextromethorphan hbr,
guaifenesin solution
DeMoulas Market Basket**

DM Max Mucus Relief

Drug Facts

**Active ingredients
(in each 20 mL)**

Dextromethorphan HBr 20 mg.

Guaifenesin 400 mg

Purpose

Dextromethorphan HBrCough Suppressant

GuaifenesinExpectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes
- temporarily relieves these symptoms occurring with a cold:
- cough due to minor throat and bronchial irritation

Warnings

Do not use • for children under 12 years of age • 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 lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis or emphysema.
- cough that occurs with too much phlegm (mucus)

When using this product • do not use more than directed

Stop use and ask a doctor if

• cough lasts for more than 7 days, comes back, or occurs with 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.

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Directions

- take only as recommended
- use dosage cup
- mL=milliliter
- do not take more than 6 doses in any 24-hour period

Age	Dose
Adults & children 12 years and older	20 mL every 4 hours
Children under 12 years of age	do not use

Other information

- each 20 mL contains: sodium 12 mg
- dosage cup provided
- store between 15-30°C (59-86°F)
- do not refrigerate

Inactive ingredients

anhydrous citric acid, dextrose, D&C Red #33, FD&C Red #40, flavors, glycerin, maltitol, propylene glycol, purified water, saccharin sodium, sodium benzoate, sucralose, Xanthan gum

Questions? Call weekdays from 9:30 AM to 4:30 PM EST at 1-877-798-5944

DM Max Mucus Relief - Product Label

"MORE FOR YOUR DOLLAR"

MARKET BASKET®

COMPARE TO Mucinex FAST-MAX™

DM MAX active ingredients*

Maximum Strength

DM Max

Mucus

Relief

CONGESTION & COUGH

**Dextromethorphan HBr
Guaifenesin**

**COUGH SUPPRESSANT /
EXPECTORANT**

ADULTS

For Ages

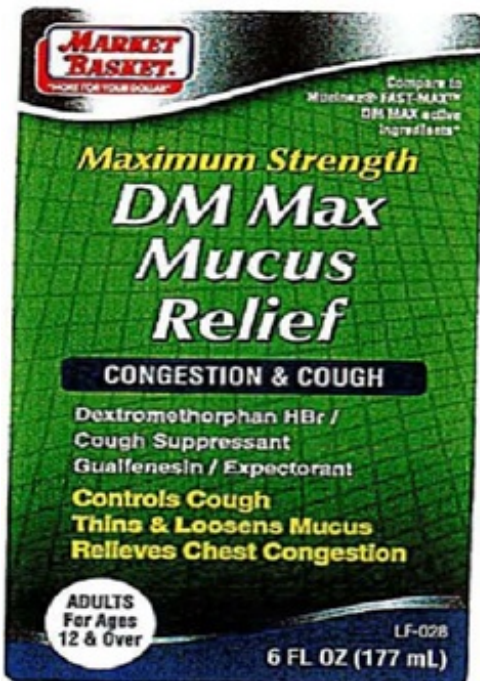
12 and Over

6 FL OZ (177 mL)

DO NOT USE IF PRINTED NECKBAND IS BROKEN OR MISSING

**DISTRIBUTED BY
DEMOULAS SUPERMARKETS INC.
875 East Street
Tewksbury, MA 01876**

*This product is not manufactured or distributed by Reckitt Benckiser Inc., distributor of Mucinex® FAST-MAX™ DM MAX.



Peel Corner to Read Complete Drug Facts and Information



DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSING
DISTRIBUTED BY: DEMICLAS SUPERMARKETS INC.
875 EAST STREET, TEWKSBURY, MA 01878



@marketbasket
Please visit us at:
thopmarketbasket.com



LF-028

Lot: Exp:

Drug Facts

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Drug Facts (continued)

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DM MAX CONGESTION AND COUGH

dextromethorphan hbr, guaifenesin solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53942-518
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)	
DEXTROSE (UNII: IY9XDZ35W2)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
MALTITOL (UNII: D65DG142WK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
SODIUM SACCHARIN (UNII: SB8ZUX40TY)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	red	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53942-518-25	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/01/2026	

Labeler - DeMoulas Market Basket (007869647)**Registrant** - AptaPharma Inc. (790523323)

Establishment

Name	Address	ID/FEI	Business Operations
AptaPharma Inc.		790523323	manufacture(53942-518)

Revised: 5/2026

DeMoulas Market Basket