MAXIMUM STRENGTH DM MAX- dextromethorphan hydrobromide, guaifenesin liquid H E B

H E B Maximum Strength DM MAX

Drug Facts

Active ingredients (in each 20 mL)

Dextromethorphan HBr 20 mg Guaifenesin 400 mg

Purpose

Dextromethorphan HBr ... Cough suppressant GuaifenesinExpectorant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes
- temporarily relieves these symploms 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 oxiclase
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

Slop 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.

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 waler, saccharin sodium, sodium benzoate, sucralose, Xanthan gum

Questions? Call weekdays from 9:30 AM to 4:30 PM EST at

1-877-798-5944

Product Label

Compare to Mucinex® FAST-MAX™

DM MAX active ingredients*

NDC 37808-518-25

H-E-B_®

Maximum Strength DM MAX

Dextromethorphan HBr / Cough Suppressant Guaifenesin / Expectorant

Congestion & Cough

Controls Cough
Thins & Loosens Mucus
Relieves Chest Congestion

Adults

For Ages 12 & Over

6 FL OZ (177mL)

LF 021

Peel Corner to Read Complete Drug Facts and Information →

DO NOT USE IF PRINTED SEAL

UNDER CAP IS TORN OR MISSING

H-E-B_®

100% GUARANTEE promise

If you aren't completely pleased with this product, we'll be happy to replace it or refund your money. You have our word on it.

0 41220 83197 5

Lot:

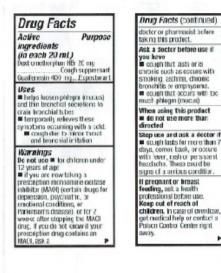
Exp:

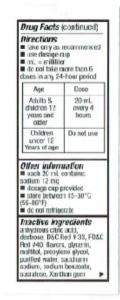
*This product is not manufactured or distributed by Reekitt Benckiser Inc., distributor of Mucinex® FAST-MAXTM DM MAX

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dextromethorphan hydrobromide, guaifenesin liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-518

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: 059QF0KO0R)		
SACCHARIN SODIUM (UNII: SB8ZUX40TY)		
SODIUM BENZOATE (UNII: OJ245FE5EU)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		
XANTHAN GUM (UNII: TTV12P4NEE)		

Product Characteristics			
Color	red	Score	
Shape		Size	
Flavor	CHERRY	Imprint Code	
Contains			

Packaging				
# Item Code Package Description		Marketing Start Date	Marketing End Date	
1	NDC:37808-518- 25	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/14/2012	

Marketing Information			
Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date

Labeler - H E B (007924756)

Registrant - AptaPharma Inc. (790523323)

Establishment			
Name	Address	ID/FEI	Business Operations
AptaPharma Inc.		790523323	manufacture(37808-518)

Revised: 12/2023 H E B