H E B MAX SEVERE CONGESTION AND COUGH MAXIUM STRENGTH- dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid
H E B

Drug Facts

Active ingredients (in each 20 mL)

Dextromethorphan HBr 20 mg
Guaifenesin 400 mg
Phenylephrine HCL 10 mg

Purpose

Dextromethorphan HBr ... Cough suppressant
Guaifenesin .....Expectorant
Phenylephrine HCL .....Nasal Decongestant

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
  • nasal congestion

Warnings

Do not use • 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.
• for children under 12 years of age

Ask a doctor before use if you have

• heart disease
• thyroid disease
• high blood pressure
• diabetes

• trouble urinating due to an enlarged prostate gland
• 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
• nervousness, dizziness or sleeplessness occur
• symptoms do not get better within 7 days or are accompanied by a fever
• cough 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 10 mg
- dosage cup provided
- store between 15-30° C (59-86° F)
- do not refrigerate

Inactive ingredients
anhydrous cilric acid, edetate disodium, FD&C Blue #1, FD&C Red #40. flavors. glycerin, propylene glycol, propyl gallate, purified water, sodium benzoate, sorbitol, 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™

Severe Congestion and Cough* active ingredients

NDC 37808-317-25

H-E-B®

Maximum Strength

**Max Severe
Congestion &
Cough
Dextromethorphan HBr /
Cough Suppressant
Guaifenesin / Expectorant
Phenylephrine HCl /
Nasal Decongestant**

Congestion & Cough

Relief of:
• Cough
• Thins & Loosens Mucus
• Nasal & Chest Congestion

Adults

For Ages 12 & Over

6 FL OZ (177mL)

LF 001 Rev.01

MADE WITH PRIDE AND CARE FOR H-E-B®

SAN ANTONIO, TEXAS, 782044

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.

LB·001 Rev.01
1208
8581-1711

0 41220 38704 5

Lot:
Exp:

***This product is not manufactured or distributed by Reckill Benckiser Inc.. distributor of Mucinex® FAST-MAX™ Severe Congeslion &Cough**

MADE WITH PRIDE & CARE FOR H·E-B®
SAN ANTONIO. TX 78204

res

H E B MAX SEVERE CONGESTION AND COUGH MAXIUM STRENGTH

dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-317
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

PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 20 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-317-25	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/14/2012	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/14/2012	

Labeler - H E B (007924756)

Registrant - AptaPharma Inc. (790523323)

Establishment

Name	Address	ID/FEI	Business Operations
AptaPharma Inc.		790523323	manufacture(37808-317)

Revised: 12/2023 H E B