

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

Discount Drug Mart

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active ingredients (in each 20 mL)

Dextromethorphan HBr, USP 20 mg

Guaifenesin, USP 400 mg

Phenylephrine HCL, USP 10 mg

Purpose

Cough Suppressant

Expectorant

Naasal Decongestant

Keep out of reach of children.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Uses

- help 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
- nasal congestion

Warnings

DO NOT USE IF PRINTED SEAL UNDER CAP IS TORN OR MISSING.

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

- 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.

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 and 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 citric acid, edetate disodium, FD and C Blue # 1, FD and Red # 40, glycerin, 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

DISCOUNT drug mart FOOD FAIR MAXIUM STRENGTH SEVERE CONGESTION & COUGH MAX product label

*COMPARE TO THE ACTIVE INGREDIENTS IN MUCINEX® FAST-MAX™ SEVERE CONGESTION AND COUGH

LF-011

DISCOUNT

drug mart

FOOD FAIR

MAXIUM STRENGTH SEVERE CONGESTION & COUGH MAX

Dextromethorphan HBr/ Guaifenesin / Phenylephrine HCL

COUGH SUPPRESSANT/

EXPECTORANT/

NASAL DECONGESTANT

Controls Cough

Thins & Loosens mucus

Relieves Nasal & Chest Congestion

Adults

For Ages 12 & Over

6 FL OZ (177 mL)

* This product is not manufactured or distributed by Reckitt Benckiser Inc., distributor of MUCINEX® FAST-MAX™ SEVERE CONGESTION AND COUGH

LB - 011 LOT: EXP:

Peel Corner to Read Complete Drug Facts and Information

SATISFACTION GUARANTEED

IF DISSASIFIED , RETURN UNUSED PORTION AND PACKAGE TO STORE WHERE PURCHASED. IF UNABLE TO RETURN TO STORE, SEND REASON FOR DISSATISFACTION, NAME, ADDRESS AND EMPTY PACKAGE TO: DISCOUNT DRUG MART, 211 COMMERCE DRIVE, MEDINA, OHIO 44256



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MUCINEX® FAST-MAX™
SEVERE CONGESTION & COUGH
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Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:53943-517

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: 059QF0K00R)			
	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:53943-517-25	177 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/10/2013	
Marketing Information				
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	OTC monograph final	part341	12/10/2013	

Labeler - Discount Drug Mart (047741335)