MUCINEX FAST-MAX SEVERE CONGESTION AND COUGH CLEAR AND COOLdextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride solution RB Health (US) LLC

Mucinex® Fast-Max [®] Severe Congestion & Cough Clear & Cool ™

Drug Facts

Active ingredients (in each 20 mL)	Purposes
Dextromethorphan HBr 20 mg	Cough suppressant
Guaifenesin 400 mg	Expectorant
Phenylephrine HCl 10 mg	Nasal decongestant

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
 - nasal congestion due to a cold
 - temporarily helps you cough less

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

- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- persistent or chronic cough 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 occur with fever
- cough comes back, or occurs with rash or headache that lasts. 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 6 doses in any 24-hour period
- measure only with dosing cup provided
- do not use dosing cup with other products
- dose as follows or as directed by a doctor
- adults and children 12 years of age and older: 20 mL in dosing cup provided every 4 hours
- children under 12 years of age: do not use

Other information

- each 20 mL contains: sodium 9 mg
- store between 20-25°C (68-77°F)
- dosing cup provided
- do not refrigerate

Inactive ingredients

anhydrous citric acid, D&C yellow no. 10, edetate disodium, FD&C blue no. 1, flavors, glycerin, propyl gallate, propylene glycol, sodium benzoate, sodium citrate, sorbitol, sucralose, water, xanthan gum

Questions?

1-866-MUCINEX (1-866-682-4639)

You may also report side effects to this phone number.

Dist. by: Reckitt Benckiser Parsippany, NJ 07054-0224 Made in England

PRINCIPAL DISPLAY PANEL - 180 mL Bottle Label

NDC 63824-541-66

MAXIMUM STRENGTH

Mucinex® FAST-MAX®

SEVERE CONGESTION & COUGH

CLEAR & **COOL** ™

Dextromethorphan HBr – Cough Suppressant Guaifenesin – Expectorant Phenylephrine HCl – Nasal Decongestant

- Controls Cough
- Relieves Nasal & Chest Congestion
- Thins & Loosens Mucus

6 FL OZ (180mL) FOR AGES 12+



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Drug Facts (continued)

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- heart disease
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Drug Facts (continued)

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Directions

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- measure only with dosing
- cup provided do not use dosing cup with
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Other information

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- dosing cup provided do not refrigerate

PEEL CORNER TO READ COMPLETE DRUG FACTS AND INFORMATION

Tamper evident: Do not use if neckband on bottle cap is broken or missing.

PARENTS: Learn about teen medicine abuse

www.StopMedicineAbuse.org



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Drug Facts (continued)

Inactive ingredients

anhydrous citric acid, D&C yellow no.10, edetate disodium.



MUCINEX FAST-MAX SEVERE CONGESTION AND COUGH CLEAR AND COOL

dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63824-541
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)				
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)				
GLYCERIN (UNII: PDC6A3C0OX)				
PROPYL GALLATE (UNII: 8D4SNN7V92)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
SODIUM BENZOATE (UNII: OJ245FE5EU)				

SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
WATER (UNII: 059QF0KO0R)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics			
Color	green	Score	
Shape		Size	
Flavor	MENTHOL	Imprint Code	
Contains			

ı	Packaging				
	# Item Package Description			Marketing Start Date	Marketing End Date
			180 mL in 1 BOTTLE; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)	07/01/2016	09/07/2024

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	07/01/2016	09/07/2024

Labeler - RB Health (US) LLC (081049410)

Revised: 10/2023 RB Health (US) LLC