

MUCINEX FAST-MAX SEVERE CONGESTION AND COUGH- dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride tablet, coated
RB Health (US) LLC

Mucinex® Fast-Max® Severe Congestion and Cough

Drug Facts

<i>Active ingredients (in each caplet)</i>	<i>Purposes</i>
Dextromethorphan HBr 10 mg	Cough suppressant
Guaifenesin 200 mg	Expectorant
Phenylephrine HCl 5 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
 - the impulse to cough to help you get to sleep
 - nasal congestion due to a cold

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
- diabetes
- high blood pressure
- thyroid disease
- 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 fever, 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 12 caplets in any 24-hour period
- adults and children 12 years of age and over: take 2 caplets every 4 hours
- children under 12 years of age: do not use

Other information

- store at 20-25°C (68-77°F)

Inactive ingredients

croscarmellose sodium, FD&C blue no. 2 aluminum lake, FD&C red no. 40 aluminum lake, methacrylic acid-ethyl acrylate copolymer, mica, microcrystalline cellulose, polyethylene glycol 3350, polysorbate 80, polyvinyl alcohol, povidone K29/32, sodium bicarbonate, talc, titanium dioxide

Questions?

1-866-MUCINEX (1-866-682-4639)You may also report side effects to this phone number.

Dist. by: RB Health (US)
Parsippany, NJ 07054-0224

Principal Display Panel - 20 Caplet Blister Pack Carton

MAXIMUM STRENGTH
NDC 63824-193-21

Mucinex®
FAST-MAX®

SEVERE CONGESTION
& COUGH

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

✓ Controls Cough

✓ Relieves Nasal & Chest Congestion

✓ Thins & Loosens Mucus

Actual Size

20 CAPLETS

FOR AGES 12+

MAXIMUM STRENGTH



Tamper evident: Do not use if carton is damaged or if printed seal on blister is broken or missing.

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& COUGH**

NDC 63824-193-21

MAXIMUM STRENGTH



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HEALTH • HYGIENE • HOME

www.mucinex.com
Patents: www.rb.com/patents

Dist. by: RB Health (US)
Parsippany, NJ 07054-0224
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3 63824 19320 5

LOT:

EXP:

MADE IN:
3141649



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Drug Facts (continued)

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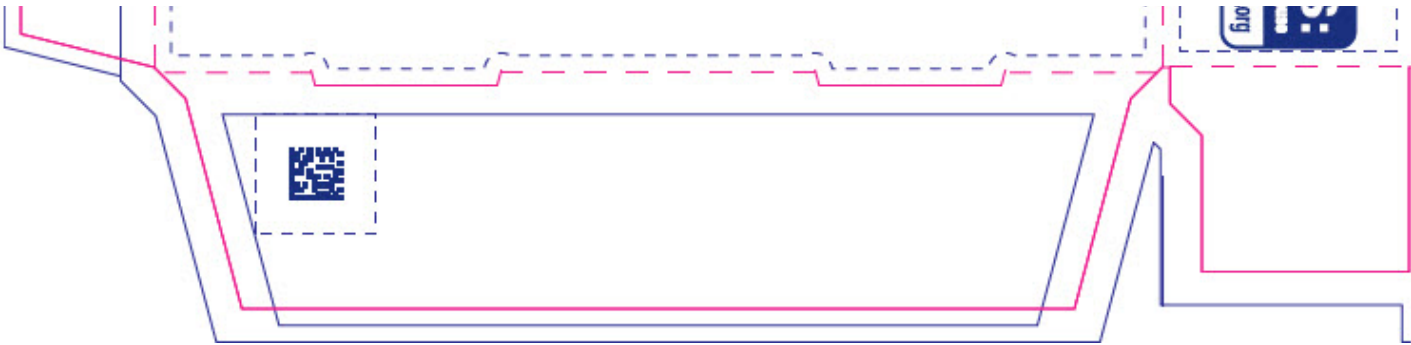


**SEVERE CONGESTION
& COUGH**

MAXIMUM STRENGTH

Maximum Strength per 4-hour dose
Do not take more than a total of 12 caplets in any 24-hour period.
Take only as directed.
Keep carton for full information.

PARENTS
Learn about team medicine at
www.StopMedicineAbuse.org



MUCINEX FAST-MAX SEVERE CONGESTION AND COUGH

dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63824-193
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients	
Ingredient Name	Strength
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
ALUMINUM OXIDE (UNII: LMI26O6933)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
MICA (UNII: V8A1AW0880)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 3350 (UNII: G2M7P15E5P)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
POVIDONE K30 (UNII: U725QWY32X)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	red	Score	no score
Shape	OVAL	Size	20mm
Flavor		Imprint Code	VV;SCC

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63824-193-20	2 in 1 CARTON	03/15/2013	05/27/2024
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:63824-193-30	3 in 1 CARTON	03/15/2013	09/01/2025
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:63824-193-21	2 in 1 CARTON	10/01/2020	
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/15/2013	

Labeler - RB Health (US) LLC (081049410)

Revised: 12/2024

RB Health (US) LLC