SEVERE CONGESTION AND COUGH- dextromethorphan hbr, guaifenesin, phenylephrine hcl solution Family Dollar Services Inc

Family Wellness 44-004

Active ingredients (in each 20 mL)

Dextromethorphan HBr 20 mg Guaifenesin 400 mg Phenylephrine HCl 10 mg

Purpose

Cough suppressant Expectorant 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

- difficulty in urination due to enlargement of the prostate gland
- thyroid disease
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- high blood pressure
- heart disease
- diabetes
- cough that occurs with too much phlegm (mucus)

When using this product

do not exceed recommended dosage.

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 persists more than 1 week, tends to recur, or is accompanied by a 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

- do not take more than directed
- do not take more than 6 doses in any 24-hour period
- mL = milliliter
- only use the dose cup provided
- adults and children 12 years and over: 20 mL in dosing cup provided every 4 hours
- children under 12 years: do not use

Other information

- each 20 mL contains: sodium 9 mg
- use by expiration date on package
- store at 25°C (77°F); excursions permitted between 15°-30°C (59°-86°F)

Inactive ingredients

anhydrous citric acid, FD&C blue #1, FD&C red #40, flavors, glycerin, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sodium metabisulfite, sorbitol, sucralose, xanthan gum

Questions or comments?

1-800-426-9391

Principal Display Panel

FAMILY Wellness™

*COMPARE TO THE ACTIVE INGREDIENTS

IN MUCINEX®
FAST-MAX® SEVERE
CONGESTION & COUGH

MAXIMUM STRENGTH
SEVERE CONGESTION
& COUGH

Dextromethorphan HBr Guaifenesin Phenylephrine HCI

Cough Suppressant Expectorant Nasal Decongestant

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

AGES
12 YEARS
& OVER

6 FL OZ (177 mL)

NDC 55319-204-45

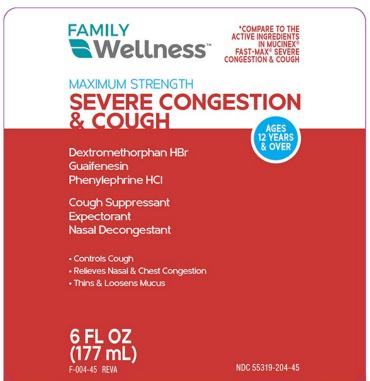
TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

PARENTS:

Learn about teen medicine abuse www.StopMedicineAbuse.org

*This product is not manufactured or distributed by RB Health (US) LLC, owner of the registered trademark Mucinex ® FAST-MAX® Severe Congestion & Cough. 50844 REV0724A00445

DISTRIBUTED BY: MIDWOOD BRANDS LLC, 500 VOLVO PKWY, CHESAPEAKE, VA 23320 USA NOT 100% SATISFIED?
Return within 30 days to the store of purchase for a refund (with receipt) or exchange.





Drug Facts (continued) 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 ■ difficulty in urination due to enlargement of the prostate gland ■ thyroid disease persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema ■ high blood pressure ■ heart disease ■ diabetes ■ cough that occurs with too much phlegm (mucus) When using this product do not exceed recommended dosage. 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 persists more than 1 week, tends to recur. or is accompanied by a fever, rash, or persistent headache. These could be signs of a serious If pregnant or breast-feeding, ask a health professional before use

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500 VOLVO PKWY, CHESAPEAKE, VA 23320 USA NOT 100% SATISFIED? Return within 30 days to the store of purchase for a refund (with receipt) or exchange

B-004-45 REVA

Family Wellness 44-004

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Control Center right away.

overdose, get medical help or contact a Poison

SEVERE CONGESTION AND COUGH

dextromethorphan hbr, guaifenesin, phenylephrine hcl solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55319-204
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)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
FD&C RED NO. 40 (UNII: WZB9127XOA)			
GLYCERIN (UNII: PDC6A3C0OX)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			
WATER (UNII: 059QF0KO0R)			
SODIUM BENZOATE (UNII: OJ245FE5EU)			
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)			
SODIUM METABISULFITE (UNII: 4VON5FNS3C)			
SORBITOL (UNII: 506T60A25R)			
SUCRALOSE (UNII: 96K6UQ3ZD4)			
XANTHAN GUM (UNII: TTV12P4NEE)			

Product Characteristics				
Color	olue	Score		
Shape		Size		
Flavor B	BERRY	Imprint Code		
Contains				

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:55319- 204-45	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/04/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	01/04/2024	

Labeler - Family Dollar Services Inc (024472631)

Establishment				
Name	Address	ID/FEI	Business Operations	
LNK International, Inc.		967626305	manufacture(55319-204), pack(55319-204)	

Revised: 1/2025 Family Dollar Services Inc