MAXIMUM STRENGTH SEVERE CONGESTION AND COUGH- dextromethorphan hbr, guaifenesin, and phenylephrine hcl liquid
SAFEWAY
Reference Label Set Id: 7f4ccd7c-9b17-428b-8cdf-42237728f9b2

Kroger Maximum Strength Severe Congestion and Cough

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
    - 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
- 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 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. (1-800-222-1222)

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
- mL = milliliter
- **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 8 mg
- low sodium
- store at room temperature
- do not refrigerate

Inactive ingredients

anhydrous citric acid, edetate disodium, FD&C Blue No. 1, FD&C Red No. 40, flavors, potassium citrate, propylene glycol, propyl gallate, purified water, sodium benzoate, sorbitol, sucralose, xanthan gum

Questions or comments?

1-866-467-2748

PRINCIPAL DISPLAY PANEL

NDC 21130-738-06

Compare to Mucinex® Fast- Max® Maximum Strength Severe Congestion & Cough Active Ingredients*

Maximum Strength

Severe Congestion & Cough
Dextromethorphan HBr - Cough Suppressant
Guaifenesin - Expectorant
Phenylephrine HCl - Nasal Decongestant

- Control Cough
- Relieves Nasal and Chest Congestion
- Thins & loosens mucus

For Ages 12+

6 FL OZ (180 mL)

Tamper evident: do not use if printed seal under cap is broken or missing.

*This product is not manufactured or distributed by RB Health (US) LLC, the distributor of Mucinex® and Fast-Max® Maximum Strength Severe Congestion & Cough.

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MAXIMUM STRENGTH SEVERE CONGESTION AND COUGH

dextromethorphan hbr, guaifenesin, and phenylephrine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21130-738
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)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
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:21130-738-06	180 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/30/2020	

Labeler - SAFEWAY (009137209)

Revised: 11/2024 SAFEWAY