

**KROGER MAXIMUM STRENGTH SEVERE CONGESTION AND COUGH-  
dextromethorphan hbr, guaifenesin, and phenylephrine hcl liquid  
KROGER COMPANY**

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**Kroger Maximum Strength Severe Congestion and Cough**

***Drug Facts***

<b><i>Active ingredients (in each 20 mL)</i></b>	<b><i>Purposes</i></b>
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 30142-738-06

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<sup>®</sup> and Fast-Max<sup>®</sup> Maximum Strength Severe Congestion & Cough.

**Distributed by:**



## KROGER MAXIMUM STRENGTH SEVERE CONGESTION AND COUGH

dextromethorphan hbr, guaifenesin, and phenylephrine hcl liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:30142-738
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RT19KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL

<b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 20 mL
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### Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>POTASSIUM CITRATE</b> (UNII: EE90ONI6FF)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>PROPYL GALLATE</b> (UNII: 8D4SNN7V92)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-738-06	180 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/03/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	09/03/2020	

**Labeler** - KROGER COMPANY (006999528)

Revised: 7/2024

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