CVS MAXIMUM STRENGTH SEVERE CONGESTION AND COUGH-dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride liquid CVS PHARMACY, INC

Reference Label Set Id: 7f4ccd7c-9b17-428b-8cdf-42237728f9b2

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

CVS Maximum Strength Severe Congestion and Cough Cooling Menthol Flavor 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 at (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 6 mg
- very low sodium
- store at room temperature
- do not refrigerate
- dosing cup provided

Inactive Ingredients

anhydrous citric acid, D&C Yellow No. 10, edetate disodium, FD&C Blue No. 1, flavor, 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# 69842-767-06

Compare to Mucinex $^{\rm B}$ Fast-Max $^{\rm B}$ Maximum Strength Severe Congestion & Cough Clear and Cool $^{\rm B}$ Active Ingredients*

Maximum Strength

Severe Congestion & Cough

Dextromethorphan HBr Cough Suppressant **Guaifenesin** Expectorant **Phenylephrine HCI** Nasal Decongestant

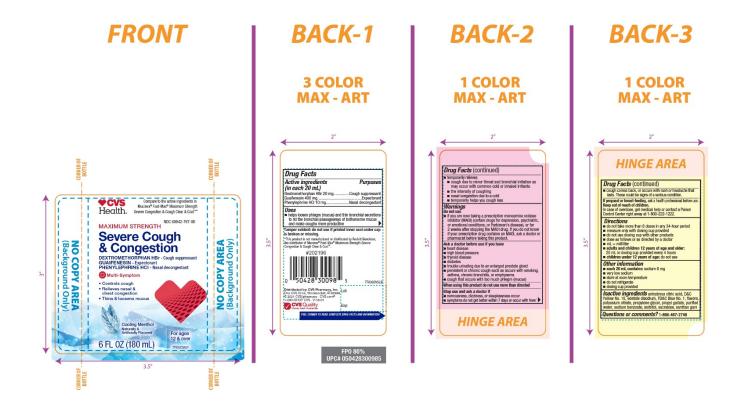
Cooling Menthol Flavor

Naturally and Artificially Flavored 6 FL OZ (180 mL)

For Ages 12+

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

*This product is not manufactured or distributed by Reckitt Benckiser, the distributor of Mucinex® Fast -Max® Maximum Strength Severe Congestion & Cough Clear & Cool™.



dextromethorphan hydrobromide, guaifenesin, and phenylephrine hydrochloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69842-767
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)	guaifenes in	400 mg in 20 mL	
<pre>phenylephrine hydrochloride (UNII: 04JA59TNSJ) (phenylephrine - UNII:1WS297W6MV)</pre>	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)		
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)		

Product Characteristics			
Color	GREEN	Score	
Shape		Size	
Flavor	MENTHOL (Cooling Menthol)	Imprint Code	
Contains			

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:69842- 767-06	180 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/31/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	03/31/2020	

Labeler - CVS PHARMACY, INC (062312574)

Revised: 9/2023 CVS PHARMACY, INC