

**CVS MUCUS RELIEF DM AND OVERNIGHT COLD AND FLU- dextromethorphan hbr, guaifenesin, acetaminophen and triprolidine hcl**  
**CVS**

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**Maximum Strength Mucus Relief DM and Overnight Cold & Flu Value Pack**

**Drug Facts**

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**Active ingredients (in each Purposes 20 mL)**

**Maximum Strength Mucus Relief DM Max**

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Dextromethorphan HBr 20 mg	Cough suppressant
Guaifenesin 400 mg	Expectorant

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**Active ingredients (in each Purposes 20 mL)**

**Nighttime Cold & Flu**

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<b>Acetaminophen 650 mg</b>	<b>Pain reliever/fever reducer</b>
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Dextromethorphan HBr 20 mg	Cough suppressant
Tripolidine HCl 2.5 mg	Antihistamine

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**Uses**

**MAXIMUM STRENGTH MUCUS RELIEF DM**

- 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

**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**

- 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***

- cough lasts more than 7 days comes back or occurs with fever, rash or headache that lasts. These could be signs of a serious condition.

### **If pregnancy 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 Centre 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 and older:** 20 mL 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
- dosing cup provided

### **Inactive ingredients (Maximum strength mucus relief DM)**

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

### **Uses (Nighttime Cold and Flu)**

- temporarily relieves these common cold and flu symptoms:
- cough
- minor aches and pains
- sore throat
- headache
- runny nose
- sneezing
- itching of the nose or throat
- itchy, watery eyes due to hay fever

- temporarily reduces fever
- controls cough to help you get to sleep

## Warnings

**Liver warnings:**This product contains acetaminophen. Severe liver damage may occur if you take\

- more than 4000 mg in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks daily while using this product

## If pregnant or breast feeding

Ask a health professional before use

## Keep Out of Reach of Children

**Overdose warning:**Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Centre right away at 1-800-222-1222.

Quick medical attention is critical for adults as well as for children, even if you do not notice any signs

## Directions

- **do not take more than directed (see overdose warnings)**
- do not take more than 4 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
- **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 10 mg
- low sodium
- store at room temperature
- do not refrigerate

## Inactive ingredients (Overnight Cold & Flu)

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

## Questions or comments?

**(1-866-467-2748)**

## **PRINCIPAL DISPLAY PANEL - Kit Carton**

### **VALUE PACK**

NDC 69842-696-12

**Compare to the active ingredients Maximum Strength Mucinex® Fast Max® DM Max\***

#### **Mucus Relief DM**

Dextromethorphan HBr • Cough Suppressant  
Guaifenesin • Expectorant  
MAXIMUM STRENGTH

MULTI-SYMPTOM

- **Controls Cough**
- **Relieves chest congestion**
- **Thins & loosens mucus**
- **4 hour dosing**

For Ages 12+

\*This product is not manufactured or distributed by Reckitt Benckiser, the distributor of Maximum Strength Mucinex® Fast-Max® DM Max.

**Compare to Mucinex® Nightshift Cold & Flu Active Ingredients\*\***

#### **Overnight Cold & Flu**

**ACETAMINOPHEN** • PAIN RELIEVER/FEVER REDUCER  
**DEXTROMETHORPHAN HBR** • COUGH SUPPRESSANT

**TRIPROLIDINE HCL** • ANTIHISTAMINE

#### **Night Time**

#### **Relief for Better Morning**

Maximum Strength per 4-hour dose

- Cough
- Fever
- Sore Throat
- Runny Nose
- Sneezing

**For Ages 12+**

**2 - 6 FL OZ (180 mL) BOTTLES / TOTAL 12 FL OZ (360 mL)**

††These product is not manufactured or distributed by Reckitt Benckister Health, distributor of Maximum Strength Mucinex® Fast Max© DM Max & Mucinex© Nightshift Cold & Flu.

**DO NOT TAKE ADULT DM MAXIMUM & NIGHTTIME COLD & FLU LIQUIDS AT THE SAME TIME.**

**TAMPER EVIDENT: DO NOT USE IF PRINTED INNER SEAL UNDER CAP IS BROKEN OR MISSING.;**

See bottle for full labeling

Distributed By:

CVS Pharmacy, Inc.

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Woonsocket, RI 02895

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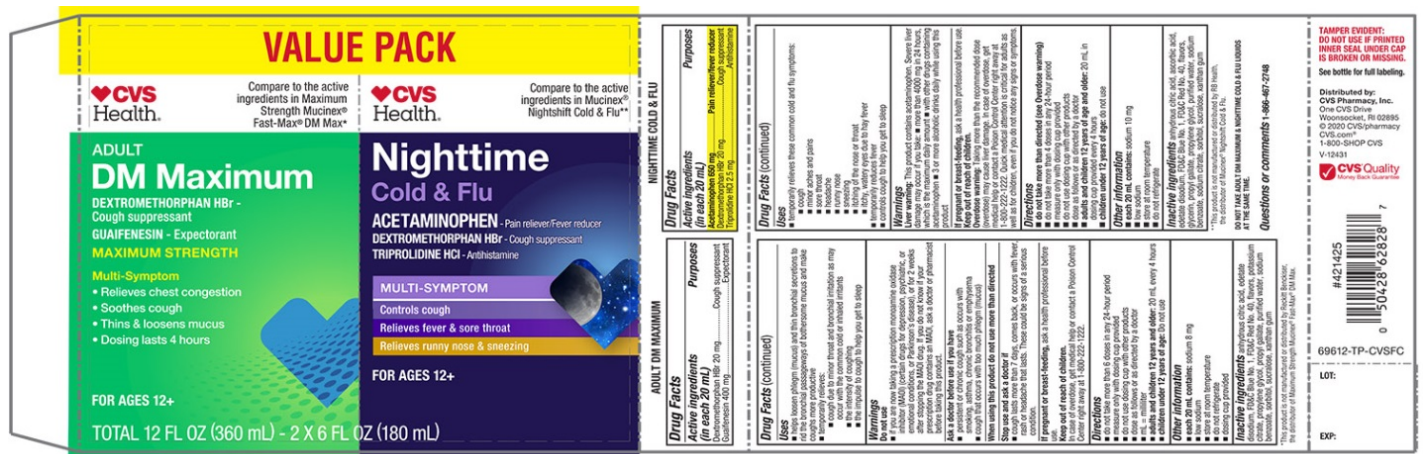
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## CVS MUCUS RELIEF DM AND OVERNIGHT COLD AND FLU

dextromethorphan hbr, guaifenesin, acetaminophen and triprolidine hcl kit

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69842-696
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### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-696-12	1 in 1 CARTON; Type 0: Not a Combination Product	03/30/2020	

## Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	180 mL
Part 2	1 BOTTLE	180 mL

## Part 1 of 2

### MAXIMUM STRENGTH MUCUS RELIEF DM

dextromethorphan hbr and guaifenesin solution

## Product Information

Route of Administration ORAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
<b>GUAIFENESIN</b> (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL

## 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)	

## Product Characteristics

Color	blue	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		180 mL in 1 BOTTLE; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/30/2020	

## Part 2 of 2

### OVERNIGHT COLD AND FLU

acetaminophen, dextromethorphan hbr and triprolidine hcl solution

## Product Information

Route of Administration ORAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg in 20 mL
<b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL
<b>TRIPROLIDINE HYDROCHLORIDE</b> (UNII: YAN7R5L890) (TRIPROLIDINE - UNII:2L8T9S52QM)	TRIPROLIDINE HYDROCHLORIDE	2.5 mg in 20 mL

## 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>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PROPYL GALLATE</b> (UNII: 8D4SNN7V92)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CITRATE, UNSPECIFIED FORM</b> (UNII: 1Q73Q2JULR)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	

## Product Characteristics

<b>Color</b>	blue	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		180 mL in 1 BOTTLE; Type 9: Other Type of Part 3 Combination Product (e.g., Drug/Device/Biological Product)		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/30/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	03/30/2020	

**Labeler** - CVS (062312574)

Revised: 8/2024

CVS