

**CVS SEVERE MULTI-SYMPATOM COUGH COLD FLU- acetaminophen,  
dextromethorphan hydrobromide, guaifenesin, phenylephrine  
hydrochloride liquid  
CVS PHARMACY**

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**CVS Adult Severe Multi-Symptom Cough Cold+Flu *Drug Facts***

***Active ingredients (in each 20 ml)***

Acetaminophen, USP 650 mg

Dextromethorphan HBr, USP 20 mg

Guaifenesin, USP 400 mg

Phenylephrine HCl, USP 10 mg

***Purposes***

Pain reliever/Fever reducer

Cough suppressant

Expectorant

Nasal decongestant

***Uses***

- temporarily relieves these symptoms occurring with a cold or flu:
- cough due to minor throat and bronchial irritation
- nasal congestion
- sinus congestion and pressure
- minor aches and pains
- sore throat
- headache
- temporarily reduces fever
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes and make coughs more productive

***Warnings***

**Liver warning**

This product contains acetaminophen. Severe liver damage may occur if you take

- more than 6 doses in any 24-hour period, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

**Allergy alert:** acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

### **Sore throat warning**

If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

### **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.
- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.

### **Ask a doctor before use if you have**

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma, chronic bronchitis, or emphysema

### **Ask a doctor or pharmacist before use if you are**

- taking the blood thinning drug warfarin
- taking any other oral nasal decongestant or stimulant
- taking any other pain reliever/fever reducer

### **When using this product do not use more than directed**

#### **Stop use and ask a doctor if**

- you get nervous, dizzy, or sleepless
- pain, cough, or nasal congestion gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- 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. Prompt medical attention is critical for adults as well as for children, even if you do not notice any signs or symptoms.

***Directions***

- do not take more than 6 doses in any 24-hour period
- do not exceed recommended dosage. Taking more than the recommended dose (overdose) may cause serious liver damage.
- measure only with dosing cup provided
- keep dosing cup with product
- ml = milliliter
- this adult product is not intended for use in children under 12 years of age

| age                                   | dose                |
|---------------------------------------|---------------------|
| adults and children 12 years and over | 20 ml every 4 hours |
| children under 12 years               | do not use          |

***Other information***

- **each 20 ml contains:** sodium 7 mg
- store at room temperature. Do not refrigerate.

***Inactive ingredients***

anhydrous citric acid, edetate disodium, FD&C red no. 40, glycerin, flavors, potassium citrate, propylene glycol, propyl gallate, purified water, sodium benzoate, sorbitol solution, sucralose, xanthan gum

***Questions or comments?***

**1-866-467-2748**

Distributed by:

**PRINCIPAL DISPLAY PANEL**

Compare to the active ingredients in Adult Robitussin Maximum Strength SEVERE Multi-Symptom Cough Cold + Flu CF MAX\*

NDC 51316-630-04

**MAXIMUM STRENGTH**

**Adult**

**SEVERE**

**Multi-Symptom**

**Cough Cold + Flu**

**ACETAMINOPHEN (Pain Reliever/Fever Reducer)**

Dextromethorphan HBr (Cough Suppressant)

Guaifenesin (Expectorant)  
Phenylephrine HCl (Nasal Decongestant)

- Cough, Sore Throat
- Body Aches, Fever
- Nasal Congestion
- Chest Congestion

Sugar-Free

No Added Alcohol

**Natural Raspberry Flavor**

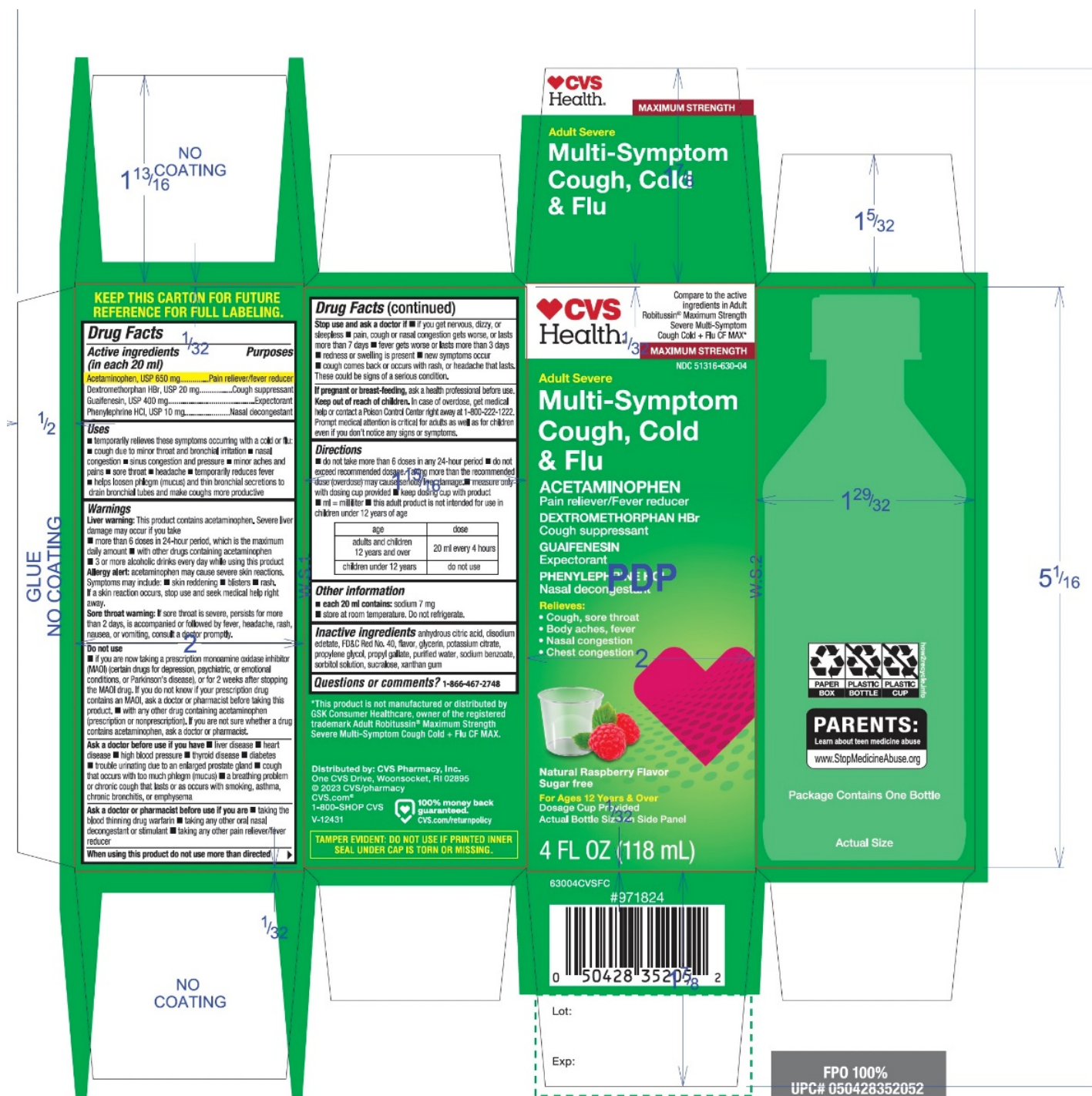
For Ages 12 & Over

8 FL OZ (237 ml)

\*This product is not manufactured or distributed by GSK Consumer Healthcare, Owner of the registered trademark Adult Robitussin Maximum Strength SEVERE Multi-Symptom Cough Cold + Flu CF MAX.



Package Label for 4FL. OZ. (118 mL)



## CVS SEVERE MULTI-SYMPTOM COUGH COLD FLU

acetaminophen, dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid

### Product Information

|                         |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:51316-630 |
| Route of Administration | ORAL           |                    |               |

| Active Ingredient/Active Moiety   |                            |   |                               |                    |
|---|----------------------------|---|-------------------------------|--------------------|
| Ingredient Name   |                            |   | Basis of Strength             | Strength           |
| ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)                    |                            |   | ACETAMINOPHEN                 | 650 mg in 20 mL    |
| 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 RED NO. 40 (UNII: WZB9127XOA)  |                            |   |                               |                    |
| GLYCERIN (UNII: PDC6A3C0OX)   |                            |   |                               |                    |
| 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   | RED                        |   | Score                         |                    |
| Shape   |                            |   | Size                          |                    |
| Flavor  | RASPBERRY, CHOCOLATE, MINT |   | Imprint Code                  |                    |
| Contains  |                            |   |                               |                    |
|   |                            |   |                               |                    |
| Packaging   |                            |   |                               |                    |
| #   | Item Code                  | Package Description                                   | Marketing Start Date          | Marketing End Date |
| 1   | NDC:51316-630-04           | 1 in 1 CARTON   | 01/09/2023                    |                    |
| 1   |                            | 118 mL in 1 BOTTLE; Type 0: Not a Combination Product |                               |                    |
| 2   | NDC:51316-630-08           | 1 in 1 CARTON   | 01/09/2023                    |                    |
| 2   |                            | 237 mL in 1 BOTTLE; Type 0: Not a Combination Product |                               |                    |
|   |                            |   |                               |                    |
| Marketing Information   |                            |   |                               |                    |
| Marketing Category  |                            | Application Number or Monograph Citation              | Marketing Start Date          | Marketing End Date |

**Labeler -** CVS PHARMACY (062312574)

Revised: 11/2023

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