

**CVS SEVERE TUSSIN CF MAX- acetaminophen, dextromethorphan hbr solution**  
**CVS PHARMACY**

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**CVS Health™ Maximum Strength Severe Tussin CF Max Drug Facts**

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

Acetaminophen, USP 650 mg

Dextromethorphan HBr, USP 20 mg

**Purposes**

Pain reliever/fever reducer

Cough suppressant

**Uses**

- temporarily relieves these symptoms occurring with a cold or flu:
- cough due to minor throat and bronchial irritation
- minor aches and pains
- sore throat
- headache
- temporarily reduces fever

**Warnings**

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

- more than 6 doses in any 24 hours 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
- cough that occurs with too much phlegm (mucus)
- a breathing problem or chronic cough that lasts or as occurs with smoking, asthma or emphysema

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

- taking the blood thinning drug warfarin
- Taking any other pain reliever/fever reducer

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

**Stop use and ask a doctor if**

- 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.**

**Overdose warning:** In case of overdose, get medical help or contact a Poison Control Center right away (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 13 mg
- store at room temperature Do not refrigerate.

**Inactive ingredients**

anhydrous citric acid, edetate disodium, FD&C red #40, glycerin, menthol, flavors, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose, xanthan gum.

**Questions or comments?**

**1-866-467-2748**

**Principal Display Panel**

Compare to active ingredients in Robitussin® Maximum Strength Severe Cough + Sore Throat\*

NDC# 69842-947-08

MAXIMUM STRENGTH

Non-Drowsy

Severe Tussin CF Max

ACETAMINOPHEN

Pain reliever / Fever reducer

DEXTROMETHORPHAN HBr

Cough Suppressant

Adult Cough & Sore Throat

Relieves:

- Cough
- Sore throat pain

For Ages 12 Years & older

Dosage cup provided

Actual Product Size on Side Panel

8 FL OZ (237 mL)

\*This product is not manufactured or distributed by Pfizer, owner of the registered trademark Robitussin Maximum Strength Severe Cough + Sore Throat.

Maximum Strength claim based on maximum levels of active ingredients per dose.

**IMPORTANT: Keep this carton for future reference on full labeling.**

Distributed by:

CVS Pharmacy, Inc.

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Woonsocket, RI 02895

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V-12431

CVS Quality

Money Back Guarantee

## CVS SEVERE TUSSIN CF MAX

acetaminophen, dextromethorphan hbr solution

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69842-947
<b>Route of Administration</b>	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

**Inactive Ingredients**

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>MENTHOL, UNSPECIFIED FORM</b> (UNII: L7T10EIP3A)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0K00R)	
<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>	PURPLE	<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	NDC:69842-947-08	1 in 1 CARTON	04/15/2019	
1		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
OTC Monograph Drug	M012	04/15/2019	

**Labeler -** CVS PHARMACY (062312574)

Revised: 2/2024

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