TUSSIN SEVERE MULTI SYMPTOM COUGH COLD PLUS FLU CF MAX- acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl solution Rite Aid Corporation

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.

Rite Aid Corporation Tussin Severe Multi-Symptom Cough Cold + Flu CF MAX 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 4,000 mg of acetaminophen in 24 hours
- 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.
- if you have ever had an allergic reaction to this product or any of its ingredients

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)
- persistent or chronic cough such 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

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.

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 5 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 15 mg
- store at 20-25°C (68-77°F). Do not refrigerate.

Inactive ingredients

anhydrous citric acid, benzyl alcohol, edetate disodium, FD&C red #40, flavor, glycerin, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol solution, sucralose, triacetin, xanthan gum

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

Compare to the active ingredients of Robitussin® Maximum Strength Severe Multi-Symptom Cough Cold + Flu CF Max

MAXIMUM STRENGTH

ADULT

tussin severe multi-symptom cough cold + flu CF MAX acetaminophen, USP 650 mg

dextromethorphan HBr, USP 20 mg
guaifenesin, USP 400 mg
phenylephrine HCl, USP 10 mg
pain reliever/fever reducer
cough suppressant
expectorant
nasal decongestant

NON-DROWSY

ALCOHOL FREE

relieves:

cough, sore throat

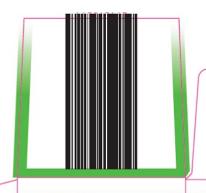
body aches, fever

nasal congestion

chest congestion

for ages 12 & over

8 FL OZ (237 mL)





MAXIMUM STRENGTH

DIA BITE



*Compare to the active ingredients of Robitus sin® Maximum Strength Severe Multi-Symptom Cough Cold + Flu CF Max

MAXIMUM STRENGTH

ADULT

tussin SeVere multi-symptom cough cold + flu

acetaminophen, USP 650 mg dextromethorphan HBr, USP 20 mg guaifenesin, USP 400 mg phenylephrine HCI, USP 10 mg

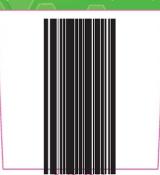
> pain reliever/fever reducer expectorant nasal decongestant

V-DROWSY LCOHOL FREE

relieves: cough, sore throat body aches, fever nasal congestion chest congestion

for ages 12 & over

8 FL 0Z (237 mL)





*Compare to the active ingredients of Robitussin® Maximum Strength Severe Multi-Symptom Cough Cold + Flu CF Max

PHARMACY MAXIMUM STRENGTH

ADULT

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> pain reliever/fever reducer expectorant nasal decongestant

NON-DROWSY ALCOHOL FREE

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8 FL 0Z (237 mL)

Drug Facts

Active ingredients (in each 20 mL)
Acetaminophen, USP 850 mg., Pah reliever/fever reducer
Dextromether phan HBr, USP 20 mg., Cough auppreaeant
Guafenesh, USP 400 mg., Expectorant
Phenylephrine HCl, USP 10 mg., Nasad decongestant

- Uses

 It emporarily relieves these symptoms occurring
 with a cold or flu:

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 Insael congestion and pressure
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 I emporarily reduces fever
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Warnings
Liver warning: This product contains acetaminophen.
Severe liver damage may occur if you take
more than 4,000 mg of acetaminophen in 24 hours

- more than 4,000 mg of a cetaminophen in 24 hours with other drugs containing a cetaminophen in 8 or more alcoholic drinks every day while using this product. Allergy slert: Acetaminophen may cause severe skin reactions. Symptoms may include: believe in the containing in

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 prescription drug contains an MAOI, ask a doctor
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 with any other drug containing acetaminophen
 (prescription or nonprescription). If you are not
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 ask a doctor or pharmacist.
 If you have ever had an allergic reaction to this
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 Ask a doctor before use if two have.

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 persistent or chronic cough such as occurs with
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- Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin

When using this product do not use more than directed

Drug Facts (continued)

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 mil.milliter
 like day in product a not intended for use
 in children under 12 years of age
 dose

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

Other information

■ each 20 mL contains: sodium 15 mg ■ store at 20-25°C (88-77°F). Do not refriger

Inactive ingredients anhydrous citric acid, berzyl alcohol, edetate disodium, FD&C red #40, flavor, glycerin, polyethylene glycol, propylene glycol purified water, sodium berzoate, sodium citrate, scribitol solution, sucralcee, triacetin, xarthan gum

Questions or comments? 1-800-719-9260

DISTRIBUTED BY: RITE AID 30 HUNTER LANE, CAMP HILL, PA 17011



Dosage Cup Included



Severe Multi-Symptom Cough Cold + Flu CF Max



LOT NO.

FXP

17Cl34 83 Cl

TUSSIN SEVERE MULTI SYMPTOM COUGH COLD PLUS FLU CF MAX

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl solution

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:11822-0086

Route of Administration

ORAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ACETAMINO PHEN (UNII: 36209 ITL9 D) (ACETAMINO PHEN - UNII: 36209 ITL9 D)	ACETAMINOPHEN	650 mg in 20 mL	
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9 D2RTI9 KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg in 20 mL	
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	400 mg in 20 mL	
PHENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	10 mg in 20 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ANHYDRO US CITRIC ACID (UNII: XF417D3PSL)		
BENZYL ALCOHOL (UNII: LKG8494WBH)		
EDETATE DISO DIUM (UNII: 7FLD9 1C8 6 K)		
FD&C RED NO. 40 (UNII: WZB9127XOA)		
GLYCERIN (UNII: PDC6 A3C0 OX)		
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
WATER (UNII: 059QF0KO0R)		
SO DIUM BENZO ATE (UNII: OJ245FE5EU)		
SO DIUM CITRATE (UNII: 1Q73Q2JULR)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		
TRIACETIN (UNII: XHX3C3X673)		
XANTHAN GUM (UNII: TTV12P4NEE)		
SORBITOL (UNII: 506T60A25R)		

Product Characteristics			
Color	RED	Score	
Shape		Size	
Flavor	BERRY	Imprint Code	
Contains			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:11822-0086-1	1 in 1 CARTON	09/22/2017	
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 final	part341	09/22/2017	

Revised: 12/2019 Rite Aid Corporation