COLD, FLU AND SORE THROAT RELIEF- acetaminophen 650 mg, dextromethorphan hbr 20 mg, guaifenesin 400 mg, phenylephrine hcl 10 mg liquid

BETTER LIVING BRANDS, LLC

Albertsons Acetaminophen 650 mg, Dextromethorphan HBr 20 mg, Guaifenesin 400 mg, Phenylephrine HCl 10 mg Liquid

Drug Facts

Active ingredients (in each 20 mL)

Acetaminophen 650 mg

Dextromethorphan HBr 20 mg

Guaifenesin 400 mg

Phenylephrine HCl 10 mg

Purposes

Pain reliever/fever reducer

Cough suppressant

Expectorant

Nasal decongestant

Uses

- temporarily relieves these common cold and flu symptoms:
- cough
- nasal congestion
- minor aches and pains
- sore throat
- headache
- stuffy nose
- sinus congestion and pressure
- temporarily reduces fever
- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus 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 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks daily 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

- 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 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

- liver disease
- heart disease
- diabetes
- high blood pressure
- thyroid disease
- 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)

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

- nervousness, dizziness, or sleeplessness occur
- pain, nasal congestion, or cough 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 fever, 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: Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Quick 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 directed (see Overdose warning)
- 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
- adults and children 12 years of age and over: 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 15 mg
- store at 20-25°C (68-77°F)
- do not refrigerate

Inactive ingredients

anhydrous citric acid, edetate disodium, FD&C blue 1, FD&C red 40, flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate dihydrate, sorbitol, sucralose, xanthan gum

Questions or comments? Call **1-877-290-4008**

COMPARE TO

Maximum Strength Mucinex® Fast-Max® Cold, Flu & Sore Throat Liquid Active Ingredients*

NDC 21130-909-03



COLD, FLU & SORE THROAT RELIEF

MAXIMUM STRENGTH

FOR AGES 12 AND OVER

ACETAMINOPHEN 650 mg

- Pain Reliever/Fever Reducer
 DEXTROMETHORPHAN HBr 20 mg
- Cough Suppressant GUAIFENESIN 400 mg
- Expectorant PHENYLEPHRINE HCI 10 mg
- Nasal Decongestant

ALL IN ONE**

Headache, Sore Throat, Sinus Pressure, Sinus Congestion, Body Pain,

Fever, Cough, Nasal Congestion

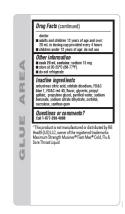
6 FL OZ (177 mL)











COLD, FLU AND SORE THROAT RELIEF

acetaminophen 650 mg, dextromethorphan hbr 20 mg, guaifenesin 400 mg, phenylephrine hcl 10 mg liquid

| Product Information | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:21130-909 |
| Route of Administration | ORAL | | |

| Active Ingredient/Active Moiety | | | | |
|--|----------------------------------|--------------------|--|--|
| Ingredient Name | Basis of Strength | Strength | | |
| ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) | 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 | | |
|----------------------|--|--|
| Strength | | |
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| Product Characteristics | | | |
|-------------------------|-------|--------------|--|
| Color | blue | Score | |
| Shape | | Size | |
| Flavor | FRUIT | Imprint Code | |
| Contains | | | |

| l | Packaging | | | | |
|---|-----------|----------------------|---|-------------------------|-----------------------|
| | # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| | | NDC:21130-909- 03 | 177 mL in 1 BOTTLE; Type 0: Not a Combination Product | 06/01/2025 | |

| Marketing Information | | | | |
|-----------------------|---|-------------------------|-----------------------|--|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date | |
| OTC Monograph Drug | M012 | 06/01/2025 | | |
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Labeler - BETTER LIVING BRANDS, LLC (009137209)

Registrant - TIME CAP LABORATORIES, INC (037052099)

| Establishment | | | | | |
|-------------------------|---------|-----------|------------------------|--|--|
| Name | Address | ID/FEI | Business Operations | | |
| MARKSANS PHARMA LIMITED | | 677604129 | manufacture(21130-909) | | |

Revised: 5/2025 BETTER LIVING BRANDS, LLC