

**DAYTIME COLD AND FLU RELIEF- acetaminophen, dextromethorphan hbr,
phenylephrine hcl capsule
BETTER LIVING BRANDS LLC.**

**693T Albertsons 21130-900 Acetaminophen, Dextromethorphan HBr and
Phenylephrine HCl softgels - Daytime Cold & Flu Softgels**

Drug Facts

Active ingredients (in each softgel)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Phenylephrine HCl 5 mg

Purpose

Pain reliever/Fever reducer

Cough suppressant

Nasal decongestant

Uses

- temporarily relieves common cold/flu symptoms:
- nasal congestion
- cough due to minor throat & bronchial irritation
- sore throat
- headache
- minor aches & pains
- fever

Warnings

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

- more than 4 doses in 24 hrs, which is the maximum daily amount for this product
- 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, lasts 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
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- persistent or chronic cough as occurs with smoking, asthma, 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, nasal congestion or cough get worse or last 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 (1-800-222-1222) right away. Quick medical attention is critical for adults & for children even if you do not notice any signs or symptoms

Directions

- take only as directed
- do not exceed 4 doses per 24 hrs
- adults & children 12 yrs & over: 2 softgels with water every 4 hrs
- children 4 to under 12 yrs: ask a doctor
- children under 4 yrs: do not use

Other information

- store at room temperature

Inactive ingredients FD&C red #40, FD&C yellow #6, gelatin, glycerin, polyethylene glycol, povidone, propylene glycol, purified water, shellac, sorbitol sorbitan solution,

titanium dioxide

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

SAFETY SEALED: DO NOT USE IF THE INDIVIDUAL
BLISTER UNIT IS OPEN OR TORN

READ AND KEEP CARTON FOR COMPLETE
WARNINGS AND PRODUCT INFORMATION

Questions? 1-877-290-4008

This product is not manufactured or distributed by Vicks VapoRub. It is a separate product. Please do not use Vicks VapoRub treatments where Vicks VapoRub and LiqueCaps are used together.

DISTRIBUTED BY: VICKS, INC./SIGNATURE CARE, LLC
P.O. BOX 1000, WASHINGTON, VA 22180-0000
1-888-273-0000

6837 0003 MADE IN INDIA

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Other Information

store at room temperature

Inactive Ingredients: FD&C Yellow #6, gelatin, glycerin, polyethylene glycol, polyethylene glycol, purified water, silica, sorbitol sodium solution, titanium dioxide.

Compare to Vicks® DayQuil® Cold & Flu LiquiCaps® active ingredients

NDC 21130-900-24



Quality Guaranteed

Daytime Cold & Flu Relief

ACETAMINOPHEN 325 mg - Pain reliever/fever reducer
DEXTROMETHORPHAN HBr 10 mg - Cough Suppressant
PHENYLEPHRINE HCl 5 mg - Nasal Decongestant

Actual Size



24 SOFTGELS*
(1 capsule of 24 softgels)

Lot #: _____

Exp Date: _____

Print/Varnish Omit Area

Varnish Omit Area

PARENTS:
Learn about teen medicine abuse
www.StopMedicineAbuse.org

RD 23008

5994

21130 76334 5

Print/Varnish Omit Area

DAYTIME COLD AND FLU RELIEF

acetaminophen, dextromethorphan hbr, phenylephrine hcl capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-900
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1W5297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
SORBITOL SOLUTION (UNII: 8KW3E207O2)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POVIDONE (UNII: FZ989GH94E)	
GLYCERIN (UNII: PDC6A3C0OX)	
SHELLAC (UNII: 46N107B71O)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GELATIN (UNII: 2G86QN327L)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
WATER (UNII: 059QF0K00R)	
SORBITAN (UNII: 6O92ICV9RU)	

Product Characteristics

Color	orange (Orange transparent)	Score	no score
Shape	OVAL	Size	21mm
Flavor		Imprint Code	70
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-900-24	1 in 1 CARTON	07/15/2023	
1		12 in 1 BLISTER PACK; 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	07/15/2023	

Labeler - BETTER LIVING BRANDS LLC. (009137209)

Registrant - TIME CAP LABORATORIES, INC (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
MARKSANS PHARMA LIMITED		925822975	manufacture(21130-900)

Revised: 2/2025

BETTER LIVING BRANDS LLC.