BERKLEY AND JENSEN ACETAMINOPHEN- acetaminophen tablet BJWC

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.

BJWC Acetaminophen Drug Facts

Active ingredient (in each caplet)

Acetaminophen 500 mg

Purpose

Pain reliever/fever reducer

Uses

- temporarily relieves minor aches and pains due to:
- the common cold
- headache
- backache
- · minor pain of arthritis
- muscular aches
- toothache
- premenstrual and menstrual cramps
- temporarily reduces fever

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.

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 have ever had an allergic reaction to this product or any of its ingredients

Ask a doctor before use if you have

liver disease

Ask a doctor or pharmacist before use if you are

taking the blood thinning drug warfarin

Stop use and ask a doctor if

- pain gets worse or lasts more than 10 days
- new symptoms occur
- fever gets worse or lasts more than 3 days
- redness or swelling is present

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). 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)

adults and children 12 years and over	 take 2 caplets every 6 hours while symptoms last do not take more than 6 caplets in 24 hours, unless directed by a doctor do not use for more than 10 days unless directed by a doctor
children under 12	ask a doctor
years	

Other information

store at 20-25°C (68-77°F)

Inactive ingredients

carnauba wax, corn starch*, croscarmellose sodium*, hypromellose, polyethylene glycol, povidone, pregelatinized starch, sodium starch glycolate*, stearic acid

*may contain one or more of these ingredients

Questions or comments?

1-800-934-1204

Principal Display Panel

Compare to the active ingredient in Extra Strength Tylenol®

berkley jensen®

EXTRA STRENGTH

ACETAMINOPHEN

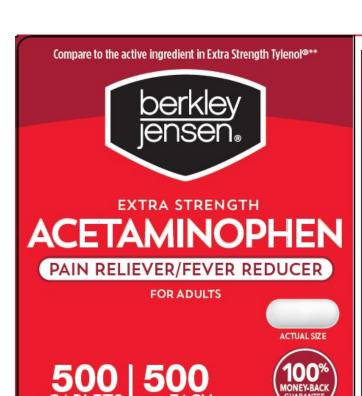
PAIN RELIEVER/FEVER REDUCER

FOR ADULTS

ACTUAL SIZE

500 CAPLETS | 500 mg EACH

100% MONEY-BACK GUARANTEE



DO NOT USE IF PRINTED SEAL UNDER CAP IS BROKEN OR MISSING

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**This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., distributor of Extra Strength Tylenol®.

GLUTEN FREE

Every Berkley Jensen product comes with our 100% Money-Back Guarantee.

Distributed by: BJ's Wholesale Club, Inc. 350 Campus Drive Marlborough, MA 01752



BERKLEY AND JENSEN ACETAMINOPHEN

acetaminophen tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68391-659

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) **ACETAMINOPHEN** 500 mg

Inactive Ingredients

Ingredient Name Strength

CARNAUBA WAX (UNII: R12CBM0EIZ)

STARCH, CORN (UNII: 08232NY3SJ)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	

Product Characteristics			
Color	WHITE	Score	no score
Shape	OVAL	Size	16mm
Flavor		Imprint Code	L484
Contains			

Pa	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:68391-659- 90	500 in 1 BOTTLE; Type 0: Not a Combination Product	03/21/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part343	03/21/2023	

Labeler - BJWC (159082692)

Revised: 3/2023 BJWC