BERKLEY AND JENSEN PAIN RELIEF EXTRA STRENGTH- 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 Pain Relief 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
- toothache
- muscular aches
- 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
- fever gets worse or lasts more than 3 days
- new symptoms occur
- 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 years	ask a doctor

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[®] SEE NEW WARNINGS EXTRA STRENGTH PAIN RELIEF ACETAMINOPHEN PAIN RELIEVER / FEVER REDUCER FOR ADULTS GLUTEN FREE 500 CAPLETS 500 mg EACH ACTUAL SIZE



BERKLEY AND JENSEN PAIN RELIEF EXTRA STRENGTH

acetaminophen tablet

Product Information									
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:68391-484					
Route of Administration	ORAL								
Active Ingredient/Active Moiety									
Ingredient Name			Basis of	Strength	Strength				
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)			ACETAMIN	OPHEN	500 mg				
Inactive Ingredients									

	Strength						
CARNAUBA WAX (UN							
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)							
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)							
PO VIDONE, UNSPECIFIED (UNII: FZ989GH94E)							
STEARIC ACID (UNII: 4ELV7Z65AP)							
STARCH, CORN (UNII: 08232NY3SJ)							
CROSCARMELLOSE SODIUM (UNII: M28 OL 1 HH48)							
Product Characteristics							
Color	WHITE	Score		no score			
Shape	CAPSULE (caplet)	Size		16 mm			
Flavor		Imprint Code		L484			
Contains							
Packaging							
# Item Code	Package Description		Marketing Start Date	Marketing End Date			
1 NDC:68391-484-90	00 in 1 BOTTLE; Type 0: Not a Combination Product		09/18/2006				
Marketing Information							
Marketing Category Application Number or Monograph Cita		itation	Marketing Start Date	Marketing End Date			
OTC monograph not final part343			09/18/2006				

Labeler - BJWC (159082692)

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BJWC