BERKLEY AND JENSEN ALLERGY RELIEF- cetirizine hydrochloride tablet, film coated
BJWC

BJWC Allergy Relief Drug Facts

Active ingredient (in each tablet)

Cetirizine HCl 10 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

Warnings

Do not use

if you have ever had an allergic reaction to this product or any of its ingredients or to an antihistamine containing hydroxyzine.

Ask a doctor before use if you have

liver or kidney disease. Your doctor should determine if you need a different dose.

Ask a doctor or pharmacist before use if you are

taking tranquilizers or sedatives.

When using this product

- drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding:

- if breast-feeding: not recommended
- if pregnant: 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 right away. (1-800-222-1222)

Directions

adults and children 6 years and over	one 10 mg tablet once daily; do not take more than one 10 mg tablet in 24 hours. A 5 mg product may be appropriate for less severe symptoms.
adults 65 years and over	ask a doctor
children under 6 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- store between 20-25°C (68-77°F)
- do not use if printed foil under cap is broken or missing

Inactive ingredients

corn starch, FD&C blue no. 1 aluminum lake, hypromellose, lactose monohydrate, magnesium stearate, polydextrose, polyethylene glycol, povidone, titanium dioxide, triacetin

Questions or comments?

1-800-934-1204

Principal Display Panel

Compare to the active ingredient in Zyrtec®

berkley Jensen®

ORIGINAL PRESCRIPTION STRENGTH

ALLERGY RELIEF

CETIRIZINE HYDROCHLORIDE TABLETS, 10mg

ANTIHISTAMINE

INDOOR & OUTDOOR ALLERGIES

24 HOUR RELIEF OF:

- SNEEZING
- RUNNY NOSE
- ITCHY, WATERY EYES
- ITCHY THROAT OR NOSE

ACTUAL SIZE

100% MONEY-BACK GUARANTEE

365 TABLETS 10 mg EACH

Compare to the active ingredient in Zyrtec®*

berkley jensen®

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PRESCRIPTION
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*This product is not manufactured or distributed by McNeil Consumer Healthcare, Division of McNeil-PPC Inc., distributor of Zyrtec®.

GLUTEN FREE

100% MONEY-BACK GUARANTEE

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

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

8 88670 01812 5

BJ #31445 Manf. #075798098

4H288 D3 S5

BERKLEY AND JENSEN ALLERGY RELIEF

cetirizine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68391-500
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	WHITE	**Score**	no score
Shape	OVAL	**Size**	10mm
Flavor		**Imprint Code**	4H2
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68391-500-87	1 in 1 PACKAGE	11/26/2013	08/01/2015
1		300 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:68391-500-88	1 in 1 PACKAGE	11/26/2013	
2		365 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078336	11/26/2013	

Labeler - BJWC (159082692)

Revised: 9/2023 BJWC