BERKLEY AND JENSEN ALLERGY RELIEF- cetirizine hydrochloride tablet 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	one 10 mg tablet once daily; do not take more than one 10 mg tablet in 24
years and over	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	ask a doctor
age	
consumers with liver or	ask a doctor
kidney disease	

Other information

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

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®

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

365 TABLETS

10mg EACH

GLUTEN FREE

Compare to the active ingredient in Zyrtec

berkley jensen.

Original Prescription Strength

ALLERGY RELIEF

CETIRIZINE HYDROCHLORIDE TABLETS, 10mg

ANTIHISTAMINE

INDOOR & OUTDOOR ALLERGIES



RELIEF OF:

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





ACTUAL SIZE

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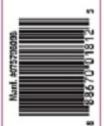
Distributed by: 84's Wholesale Club 25's Rosearch Chine Westborough, MAOLSB1 Questons? 3-800-834-1204

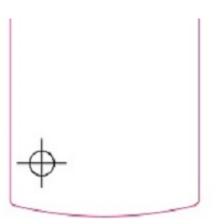
Every Berkley Jersen product connec with our 100% Money-Back Guarantee. It's just one of the many ways a fil's membership gives you MORE.





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Drug Facts

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Cetirizine HCI 10 mg......Antihistamine

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Drug Facts (continued)

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Questions or comments? 1-800-934-1204

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

BERKLEY AND JENSEN ALLERGY RELIEF

cetirizine hydrochloride tablet

Product Information

Route of Administration ORAL

Active Ingredient/Active Majety

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Ingredient Name	Basis of Strength	Strength	
CETIRIZINE HYDRO CHLO RIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	10 mg	

Inactive Ingredients				
Ingredient Name	Strength			
STARCH, CORN (UNII: O8232NY3SJ)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
LACTO SE MO NO HYDRATE (UNII: EWQ57Q8I5X)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
POLYDEXTROSE (UNII: VH2XOU12IE)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
PO VIDO NE, UNSPECIFIED (UNII: FZ989GH94E)				
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)				
TRIACETIN (UNII: XHX3C3X673)				

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

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:68391-500-87	1 in 1 CARTON	11/26/2013			
1		300 in 1 BOTTLE; Type 0: Not a Combination Product				
2	NDC:68391-500-88	1 in 1 CARTON	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: 12/2019 BJWC