

**CETIRIZINE HCL- cetirizine hcl capsule**  
**KROGER COMPANY**

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***Cetirizine HCl Capsules, 10 mg dye free-Kroger***

***Active ingredient (in each capsule)***

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
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic drinks
- be careful when driving a motor vehicle or operating machinery
- in rare cases, you may experience itching after stopping cetirizine. Consult your healthcare provider if this happens.

**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.

**Directions**

adults and children 6 years and over	one 10 mg capsule once daily; do not take more than one 10 mg capsule 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 at 20°-25°C (68°-77°F)
- avoid high humidity and excessive heat above 40°C (104°F)
- protect from light

**Inactive ingredients**

gelatin, glycerin, mannitol, pharmaceutical ink, polyethylene glycol, purified water, sodium hydroxide, sorbitan, sorbitol

**Questions or comments?**

call toll free **1-888-235-2466**(Mon - Fri 9AM - 5PM EST)

\*ZYRTEC® IS A REGISTERED TRADEMARK OF KENVUE INC.

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

**KEEP THIS CARD FOR COMPLETE WARNINGS AND PRODUCT INFORMATION**

**PROUDLY DISTRIBUTED  
BY THE KROGER CO.  
CINCINNATI, OHIO 45202  
‡OUR PHARMACIST RECOMMENDED**

**Quality Guaranteed**

**Kroger.com/guarantee**

800-632-6900

L0001257

R0326

Lot No.:

Exp. Date:

**Principal Display Panel-25s count**

**NDC 59450-020-86**

**Kroger®**

**health**

**COMPARE TO THE ACTIVE**

**INGREDIENT IN ZYRTEC®†**

**Dye Free**

**Allergy Relief**

**Cetizine HCl Capsules, 10 mg**

**Antihistamine**

**Indoor & Outdoor Allergies**

**UP TO**

**24HR**

**RELIEF**

**24 HOUR RELIEF OF:  
SNEEZING, RUNNY NOSE  
ITCHY, WATERY EYES  
ITCHY THROAT OR NOSE**

**25 SOFTGELS†**

**†LIQUID-FILLED CAPSULES**



# Dye Free Allergy Relief

UP TO  
24 HR  
RELIEF

**Cetirizine HCl Capsules, 10 mg**  
Antihistamine  
Indoor & Outdoor Allergies

ACTUAL SIZE



**24 HOUR RELIEF OF:**  
SNEEZING, RUNNY NOSE  
ITCHY, WATERY EYES  
ITCHY THROAT OR NOSE

**25 SOFTGELS<sup>†</sup>**  
**LIQUID-FILLED CAPSULES**

**Drug Facts (continued)**

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- protect from light

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gelatin, glycerin, mannitol, pharmaceutical ink, polyethylene glycol, purified water, sodium hydroxide, sorbitan, sorbitol

**Questions or comments?**  
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**Drug Facts (continued)**

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ask a doctor  
adults 65 years and over: ask a doctor  
ask a doctor  
children under 6 years of age: ask a doctor  
consumers with liver or kidney disease: ask a doctor

**Warnings**  
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.

**Drug Facts (continued)**

**When using this product**  
drowsiness may occur  
alcohol, sedatives, and tranquilizers may increase drowsiness  
avoid alcoholic drinks  
be careful when driving a motor vehicle or operating machinery. In rare cases, you may experience itching after stopping machinery. Consult your healthcare provider if this happens.  
Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.  
If pregnant or breast-feeding: not recommended

**Drug Facts**

**Active ingredient (in each capsule)**  
Cetirizine HCl 10 mg  
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

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**KEEP THIS CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION**

L0001257  
R0326  
Lot No.  
Exp. Date:

## \*Liquid-filled capsules

### CETIRIZINE HCL

cetirizine hcl capsule

#### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:59450-020
<b>Route of Administration</b>	ORAL		

#### Active Ingredient/Active Moiety

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

#### Inactive Ingredients

Ingredient Name	Strength
<b>GELATIN</b> (UNII: 2G86QN327L)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>MANNITOL</b> (UNII: 3OWL53L36A)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>SORBITAN</b> (UNII: 6O92ICV9RU)	
<b>SORBITOL</b> (UNII: 506T60A25R)	

#### Product Characteristics

<b>Color</b>	white (clear)	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	13mm
<b>Flavor</b>		<b>Imprint Code</b>	CE2
<b>Contains</b>			

#### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59450-020-86	1 in 1 CARTON	05/25/2026	
1		25 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:59450-020-15	1 in 1 CARTON	05/25/2026	
2		40 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA022429	05/25/2026	

**Labeler** - KROGER COMPANY (006999528)

**Registrant** - Bionpharma Inc. (079637826)

## Establishment

Name	Address	ID/FEI	Business Operations
Patheon Softgels Inc.		002193829	manufacture(59450-020)

Revised: 4/2026

KROGER COMPANY