

**ALL DAY ALLERGY RELIEF- cetirizine hcl capsule**  
**CHAIN DRUG MARKETING ASSOCIATION INC.**

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***Drug Facts***

***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

**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***

FD&C yellow #6, gelatin, glycerin, mannitol, pharmaceutical ink, polyethylene glycol, purified water, sodium hydroxide, sorbitan, sorbitol

***Questions or comments?***

call toll free **1-888-235-2466**

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**Distributed by C.D.M.A., Inc.©**

**43157 W 9 Mile Rd**

**Novi, MI 48375**

**www.qualitychoice.com**

**Questions: 248-449-9300**

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

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

**Principal Display Panel**

**QC <sup>®</sup>**

**QUALITY CHOICE**

**NDC 63868-433-25**

**\*Compare to the Active Ingredient in Zyrtec<sup>®</sup>**

**All Day Allergy Relief**

**Cetirizine HCl Capsules, 10 mg**

**Antihistamine**

**Indoor & Outdoor**

**Allergies**

**24 Hour Relief of:**

**Sneezing**

**Runny Nose**

**Itchy, Watery Eyes**

**Itchy Throat or Nose**

**25 softgels\*\***

**(\*\*Liquid-Filled Capsules)**

**Front**

NDC 63868-433-25



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Active Ingredient in  
Zyrtec®

# All Day Allergy Relief

Cetirizine HCl Capsules, 10 mg  
Antihistamine

**Indoor & Outdoor  
Allergies**

**24 Hour Relief of:**

Sneezing  
Runny Nose  
Itchy, Watery Eyes  
Itchy Throat or Nose



**25** Softgels\*\*  
(\*\*Liquid-Filled Capsules)

**Back**



## ALL DAY ALLERGY RELIEF

cetirizine hcl capsule

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-433
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
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (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)				
<b>Product Characteristics</b>				
<b>Color</b>	orange	<b>Score</b>	no score	
<b>Shape</b>	OVAL	<b>Size</b>	13mm	
<b>Flavor</b>		<b>Imprint Code</b>	CE1	
<b>Contains</b>				
<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:63868-433-25	1 in 1 BOX	01/25/2019	
1		25 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>		<b>Marketing Start Date</b>	<b>Marketing End Date</b>
NDA	NDA022429		01/25/2019	

**Labeler** - CHAIN DRUG MARKETING ASSOCIATION INC. (011920774)

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

## Establishment

Name	Address	ID/FEI	Business Operations
Patheon Softgels Inc.		002193829	manufacture(63868-433)

Revised: 12/2022

CHAIN DRUG MARKETING ASSOCIATION INC.