#### 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

<sup>\*</sup>All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Zyrtec  $^{\circ}$ .

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Questions: 248-449-9300

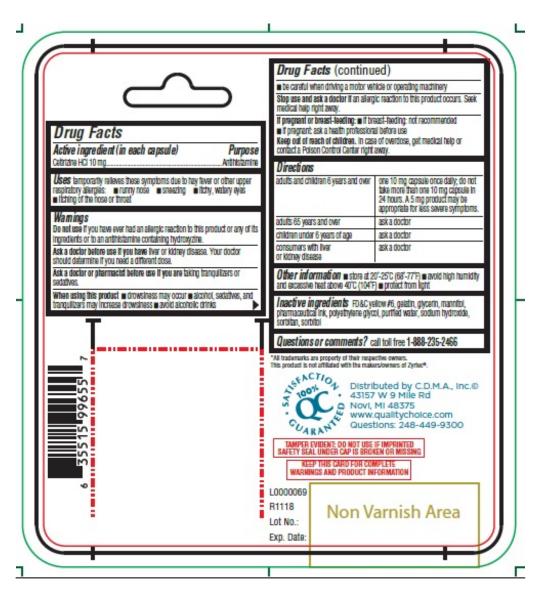
#### TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

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

**Principal Display Panel** QC<sub>®</sub> **QUALITY CHOICE** NDC 63868-433-25 \*Compare to the Active Ingredient in Zyrtec® All Day Allergy Relief Cetirizine HCI 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



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ALL DAY ALLERGY R 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 Streng					Strength	
<b>CETIRIZINE HYDROCHLORIDE</b> (UNII: 640047KTOA) (CETIRIZINE - UNII:Y07261ME24)			CETIRIZ INE HYDROCHLORIDE		10 mg	
Inactive Ingredients						
Ingredient Name					trength	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)						
FD&C TELLOW NO. 6 (UNII: H//						

MANNITOL (UNII:	30WL53L36	A)					
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)							
WATER (UNII: 059QF0K00R)							
SODIUM HYDROXIDE (UNII: 55X04QC32I)							
SORBITAN (UNII: 6092ICV9RU)							
SORBITOL (UNII:	506T60A25P	<b>(</b> )					
Product Cha	racterist	ics					
Color		orange	Score		no score		
Shape		OVAL	Size				
Flavor			Imprint Code				
Contains							
Packaging							
# Item Code		Package Description		Marketing Start Date	Marketing End Date		
<b>1</b> NDC:63868- 433-25	1 in 1 BOX	<		01/25/2019			
1 25 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product							
Marketing	ı Inforn	nation					
Marketing Marketing Category		lication Nur	nber or Monograph tation	Marketing Start Date	Marketing End Date		
Marketing		lication Nur Ci <sup>r</sup>		-	-		

Labeler - CHAIN DRUG MARKETING ASSOCIATION INC. (011920774)

# Registrant - Bionpharma Inc. (079637826)

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

Revised: 12/2022

#### CHAIN DRUG MARKETING ASSOCIATION INC.