

CETIRIZINE HCL- cetirizine hcl capsule
TARGET Corporation

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?

call **1-800-910-6874**

*All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Zyrtec[®].

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

KEEP THIS CARD FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

Dist. by Target Corp., Mpls., MN 55403

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R1019

Principal Display Panel

NDC 11673-242-15

**Compare to active ingredient in Zyrtec[®]*
allergy relief**

cetirizine HCl capsules, 10 mg antihistamine

indoor and outdoor allergies

24 hour relief of:

- sneezing
- runny nose
- itchy, watery eyes
- itchy throat or nose

24 HOUR RELIEF

ACTUAL SIZE

40 SOFTGELS **

up & upTM

40 SOFTGELS ** (**LIQUID-FILLED CAPSULES)
10 mg EACH

front

Compare to active ingredient
in Zyrtec®

NDC 11673-242-15

allergy relief

cetirizine HCl
capsules, 10 mg
antihistamine

indoor and outdoor allergies

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back

Drug Facts

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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

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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

Drug Facts (continued)

■ 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

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Lot No.:

Exp. Date:

TAMPER EVIDENT:
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COMPLETE WARNINGS
AND PRODUCT
INFORMATION**

CETIRIZINE HCL

cetirizine hcl capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-242
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

UNII:YO7261ME24)			HYDROCHLORIDE	10 mg
Inactive Ingredients				
Ingredient Name				Strength
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)				
GELATIN (UNII: 2G86QN327L)				
GLYCERIN (UNII: PDC6A3C0OX)				
MANNITOL (UNII: 3OWL53L36A)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
WATER (UNII: 059QF0KO0R)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
SORBITAN (UNII: 6O92ICV9RU)				
SORBITOL (UNII: 506T60A25R)				
Product Characteristics				
Color	orange	Score	no score	
Shape	OVAL	Size	13mm	
Flavor		Imprint Code	CE1	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-242-15	1 in 1 POUCH	10/30/2019	
1		40 in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
NDA	NDA022429		10/30/2019	

Labeler - TARGET Corporation (006961700)

Registrant - Bionpharma Inc. (079637826)

Establishment

Name	Address	ID/FEI	Business Operations
Patheon Softgels Inc.		002193829	manufacture(11673-242)