

ALLERGY RELIEF- cetirizine hcl tablet
Pioneer Life Sciences, LLC

Allergy relief

Active Ingredient (in each caplet)

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 sedatives or tranquilizers.

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 (1-800 222-1222) right away.

Directions

Adults and children 6 years and older: One 10 mg tablet once daily; do not take more than one 10 mg tablet 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 between 20°C to 25°C (68°F to 77°F)

Inactive Ingredients

Lactose, Povidone, Maize Starch, Magnesium Stearate, Opadry White, Titanium Dioxide, Hypromellose, Polyethylene Glycol

Questions or Comments?

Call 1-732-698-5070 Mon-Fri, 9am-5pm EST or visit www.gencare.health

24 Hour Relief

†compare to ZYRTEC[®] ALLERGY active ingredients.

GenCare
generic healthcare

ORIGINAL RX STRENGTH

Allergy Relief

antihistamine

Cetirizine HCl 10 mg

24 hour Relief from Indoor/Outdoor Allergies

- Sneezing, Runny Nose
- Itchy Watery Eyes
- Itchy Throat or Nose

NDC 72090-028-01

300 caplets not actual size



TAMPER EVIDENT Do not use if safety seal under cap is broken or missing.

Drug Facts

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Cetirizine HCl 10 mgAntihistamine

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(continued under label)

†This product is not manufactured or distributed by ©Johnson & Johnson Consumer Inc., owner of the Registered Trademark Zyrtec® Allergy

Distributed by: **GenCare Consumer Products, LLC** 40E Collins Ln, Suite A, East Brunswick, NJ 08816 www.genicare.health

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Lot No. Exp. Dt.

PEEL HERE

Drug Facts (continued)

Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.

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■ 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 (1-800-222-1222) right away.

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Inactive Ingredients: Lactose, Povidone, Maltose Starch, Magnesium Stearate, Opadry white, Titanium Dioxide, Hypromellose, Polyethylene Glycol.

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ALLERGY RELIEF			
cetirizine hcl tablet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72090-028
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	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDWL1A)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3W0)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
STARCH, CORN (UNII: O8232NY35J)			

POVIDONE (UNII: FZ989GH94E)

Product Characteristics

Color	white	Score	no score
Shape	BULLET (Barrel shape)	Size	8mm
Flavor		Imprint Code	CTN;10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72090-028-01	300 in 1 BOTTLE; Type 0: Not a Combination Product	10/03/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077829	10/03/2024	

Labeler - Pioneer Life Sciences, LLC (014092742)

Registrant - Pioneer Life Sciences, LLC (014092742)

Revised: 10/2024

Pioneer Life Sciences, LLC