

ALLERGY RELIEF- cetirizine hydrochloride tablet, coated
CHAIN DRUG CONSORTIUM

PRV-1194A-2019-1030

Drug Facts

Active ingredient (in each tablet)

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
- 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 right away. (1-800-222-1222)

Directions

adults and children 6 years and over	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 to 25°C (68 to 77°F)
- retain carton for complete product information and warnings

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide.

Questions or comments?

1-844-705-4384

PRINCIPAL DISPLAY PANEL

Premier Value®

COMPARE TO THE ACTIVE INGREDIENT IN ZYRTEC® ALLERGY†

2 Week Supply

24 Hour

ALLERGY RELIEF

Indoor & Outdoor Allergy Relief

Original Prescription Strength

CETIRIZINE HYDROCHLORIDE TABLETS, 10 mg

For Relief of: • Sneezing • Runny Nose

- Itchy, Watery Eyes
- Itchy Throat or Nose

Antihistamine

14 TABLETS

10 MG EACH

ACTUAL SIZE

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Drug Facts (continued)
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 If pregnant or breast-feeding:
 ■ if breast-feeding: not recommended before use;
 ■ if pregnant: ask a health professional before use.
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INK AND COATING FREE FOR LOT AND EXPIRATION STAMPING

5 52930 98607 8

F1194A01PRV_R0

DO NOT USE IF IMPRINTED SEAL UNDER CAP IS BROKEN OR MISSING

This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Zyrtec®/AllergyT.

MADE IN INDIA
 407 East Lancaster Avenue
 Wayne, PA 19087
 Pharmacy Value Alliance, LLC
 Distributed By:
 1-844-705-4384

Questions or comments?
 cellulose, polyethylene glycol, titanium dioxide monohydrate, magnesium stearate, microcrystalline cellulose, croscarmellose sodium, hypromellose, lactose dioxides, colloidal silicon dioxide

Inactive ingredients (continued)

COMPARE TO THE ACTIVE INGREDIENT IN ZYRTEC® ALLERGYT

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2 Week Supply

ALLERGY RELIEF Indoor & Outdoor Allergy Relief Original Prescription Strength

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Antihistamine

24hr

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Actual Size 

PV IMPRINTED SEAL UNDER CAP

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PV IMPRINTED SEAL UNDER CAP

NC

ALLERGY RELIEF

cetirizine hydrochloride tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-294
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)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	

Product Characteristics

Color	white (white to off white)	Score	2 pieces
Shape	RECTANGLE (rounded off rectangular)	Size	9mm
Flavor		Imprint Code	G;4
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-294-14	1 in 1 CARTON	11/01/2019	04/30/2026
1		14 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:68016-294-30	1 in 1 CARTON	11/01/2019	07/31/2026
2		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:68016-294-60	1 in 1 CARTON	11/01/2019	07/31/2026
3		60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

4	NDC:08010-294-90	1 in 1 CARTON	11/01/2019	07/31/2026
4		90 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
ANDA	ANDA209274		11/01/2019	07/31/2026

Labeler - CHAIN DRUG CONSORTIUM (101668460)

Revised: 10/2024

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