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	one 10 mg tablet once daily; do not take more than one 10 mg tablet in 24 hours. A
years and over	5 mg product may be appropriate for less severe symptoms.
adults 65 years and	ask a doctor

over	
children under 6 years	sask a doctor
of age	
consumers with liver	ask a doctor
or kidney disease	

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

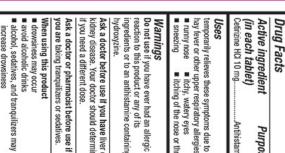
- Itchy, Watery Eyes
- Itchy Throat or Nose

Antihistamine

14 TABLETS

10 MG EACH

ACTUAL SIZE



adults and children 6 years and over

one 10 mg tablet once daily; one take more than one 10 m tablet in 24 hours. A 5 mg

Directions

overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Keep out of reach of children. In case of

before use.

be careful when driving a motor vehicle or operating machinery

> liver or kidney consumers with years of age children under 6

> > ask a doctor ask a doctor ask a doctor for less severe symptoms. product may be appropriate

 retain carton for complete product information Other information ■ store between 20-25°C (68-77°F)

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Distributed By: Pharmacy Value Alliance, LLC 407 East LancasterAvenue Vayne, PB 19087 Alexan, Mayne, PR 1004 any re

Questions or comments?

Inactive ingredients Drug Facts (continued)

cellulose, polyethylene glycol, titanium dioxide monohydrate, magnesium stearate, microcrystalline qioxiqe' croecsume pose soqinm' pilborme (poe' lactose

1-844-102-4384

Ask a doctor before use if you have liver or kidney disease. Your doctor should determine ngredients or to an antihistamine containing you need a different dose

adults 65 years and over

Warnings
Do not use if you have ever had an allergic reaction to this product or any of its hay fever or other upper respiratory allergies:

runny nose itchy, watery eyes
sneezing itching of the nose or throat

Anthistamine Purpose

■ if breast-feeding: ■ if breast-feeding: not recommended Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right pregnant ask a health professional

This product is not manufactured or distributed by McNei Consumer Healthcare, distributor of Zyrtec® Allergy. DO NOT I

EXPIRATION STAMPING FOR LOT AND 0362 0986 ∞

INK AND COATING FREE

Drug Facts (continued)

OT USE IF IMPRINTED CAP IS BROKEN OR A

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₽ op COMPARE TO THE ACTIVE INGREDIENT IN Premier ZYRTEC® ALLERGY† Value 24HR 2 Week Supply Indoor & Outdoor Allergy Relief Original Prescription Strength RELIEF CETIRIZINE HYDROCHLORIDE TABLETS, 10 mg •Sneezing • Runny Nose •Itchy, Watery Eyes For Relief of: 14 TABLETS

If for any reason you are not salisfied with this product, please return it to the store where purchased for a full refund.

(A)

COMPARE TO THE ACTIVE INGREDIENT IN Premier ZYRTEC® ALLERGY† Value 24HR 2 Week Supply Indoor & Outdoor Allergy Relief ERGY Original Prescription Strength CETIRIZINE HYDROCHLORIDE TABLETS, 10 mg •Sneezing • Runny Nose •Itchy, Watery Eyes •Itchy Throat or Nose TABLETS

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 HYDRO CHLO RIDE (UNII: 64O047KTOA) (CETIRIZINE - UNII: YO7261ME24)	CETIRIZINE HYDROCHLORIDE	10 mg	

Inactive Ingredients			
Ingredient Name	Strength		
LACTO SE MO NO HYDRATE (UNII: EWQ57Q8I5X)			
CROSCARMELLOSE SODIUM (UNII: M28 OL1HH48)			
SILICON DIO XIDE (UNII: ETJ7Z6 XBU4)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
CELLULO SE, MICRO CRYSTALLINE (UNII: OP1R32D61U)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)			
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)			

Product Characteristics			
Color	white (white to off white)	Score	2 pieces
Shape	RECTANGLE (rounded off rectangualr)	Size	9 mm
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	11/0 1/20 19	
1		14 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:68016-294- 30	1 in 1 CARTON	11/0 1/20 19	
2		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:68016-294- 60	1 in 1 CARTON	11/0 1/20 19	
3		60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:68016-294- 90	1 in 1 CARTON	11/0 1/20 19	
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	11/0 1/20 19	

Labeler - CHAIN DRUG CONSORTIUM (101668460)

Revised: 10/2019 CHAIN DRUG CONSORTIUM