

**ZYRTEC ALLERGY- cetirizine hydrochloride tablet, film coated**  
**RedPharm Drug Inc**

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**Zyrtec<sup>®</sup> Allergy**

***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)
- **do not use if clamshell is opened, or if foil inner seal imprinted with "ZYRTEC®" is broken or missing**
- meets USP Dissolution Test 2

## Inactive ingredients

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

## Questions?

call **1-800-343-7805** (toll-free) or **215-273-8755** (collect)

## PRINCIPAL DISPLAY PANEL

Original Prescription Strength

NDC 50580-726-36

ZYRTEC®

ALLERGY

Cetirizine HCl tablets

10 mg /antihistamine

Indoor & Outdoor Allergies

24

hour

Relief of

- Sneezing

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

30 Tablets  
10 mg each

(Actual Size)

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**Important: Read all product information before using. Keep this card for important information.**

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# ZYRTEC®

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10 mg / antihistamine

**ALLERGY**

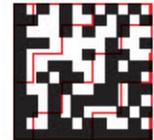
## Indoor & Outdoor Allergies

The trade dress of this ZYRTEC® package is subject to trademark protection.

Active ingredient made in Switzerland

Distributed by:  
**JOHNSON & JOHNSON CONSUMER INC.**  
McNeil Consumer Healthcare Division  
Fort Washington, PA 19034 USA |  
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USD 601,012; USD 606,856; USD 620,359;  
US 7,866,475

30040818



## ZYRTEC ALLERGY

cetirizine hydrochloride tablet, film coated

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:67296-1484(NDC:50580-726)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CETIRIZINE HYDROCHLORIDE</b> (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	10 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	

**MICROCRYSTALLINE CELLULOSE** (UNII: OP1R32D61U)

**POLYETHYLENE GLYCOL, UNSPECIFIED** (UNII: 3WJQ0SDW1A)

**TITANIUM DIOXIDE** (UNII: 15FIX9V2JP)

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	2 pieces
<b>Shape</b>	RECTANGLE (rounded-off rectangular biconvex tablet)	<b>Size</b>	9mm
<b>Flavor</b>		<b>Imprint Code</b>	ZYRTEC;10;MG
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67296-1484-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2008	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA019835	01/01/2008	

**Labeler** - RedPharm Drug Inc (828374897)

### Establishment

Name	Address	ID/FEI	Business Operations
RedPharm Drug		828374897	repack(67296-1484)

Revised: 2/2023

RedPharm Drug Inc