

**ZYRTEC ALLERGY- cetirizine hydrochloride capsule, liquid filled**  
**Johnson & Johnson Consumer Inc.**

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**ZYRTEC Allergy**

***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
- 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 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
- **do not use if clamshell is opened or if foil inner seal imprinted with “Sealed For Your Safety” is broken or missing**

## Inactive ingredients

gelatin, glycerin, mannitol, pharmaceutical ink, polyethylene glycol 400, purified water, sodium hydroxide, sorbitan, sorbitol

## Questions?

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

## PRINCIPAL DISPLAY PANEL

### ***Original Prescription Strength***

NDC 50580-786-12

**ZYRTEC<sup>®</sup>**

### ***ALLERGY***

### ***INDOOR + OUTDOOR***

### ***ALLERGIES***

***Cetirizine HCl/***

***antihistamine***

**10 mg capsules**

**LIQUID**

**GELS**

**24**

**HOUR**

**RELIEF OF**

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

**(Actual Size)**

**12**

**LIQUID GELS\***

**\*LIQUID-FILLED CAPSULES**

**10 mg each**



• Itchy Throat or Nose



(Actual Size)

**12**  
**LIQUID GELS\***

\*LIQUID-FILLED CAPSULES

**10 mg each**

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### Drug Facts

(continued)

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# ZYRTEC®

**Cetirizine HCl/  
antihistamine  
10 mg capsules**

**ALLERGY**



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

Active ingredient made in India

Distributed by:

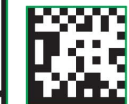
**JOHNSON & JOHNSON CONSUMER INC.**

McNeil Consumer Healthcare Division

Fort Washington, PA 19034 USA

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Pat. www.jjcipts.com



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# ZYRTEC ALLERGY

cetirizine hydrochloride capsule, liquid filled

## Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50580-786
Route of Administration	ORAL		

## Active Ingredient/Active Moiety

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

## Inactive Ingredients

Ingredient Name	Strength
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
MANNITOL (UNII: 3OWL53L36A)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SORBITAN (UNII: 6O92ICV9RU)	
SORBITOL (UNII: 506T60A25R)	

## Product Characteristics

Color	white (Clear)	Score	no score
Shape	OVAL	Size	13mm
Flavor		Imprint Code	CZ 10
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-786-12	1 in 1 PACKAGE	04/15/2024	
1		12 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:50580-786-25	1 in 1 PACKAGE	04/15/2024	
2		25 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:50580-786-40	1 in 1 PACKAGE	04/15/2024	
3		40 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:50580-786-65	65 in 1 PACKAGE, COMBINATION; Type 0: Not a Combination Product	04/15/2024	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA213105	04/15/2024	

**Labeler** - Johnson & Johnson Consumer Inc. (878046358)

Revised: 1/2024

Johnson & Johnson Consumer Inc.