

**CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet, film coated**  
**Aphena Pharma Solutions - Tennessee, LLC**

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**Cetirizine HCL Tablet 10 mg**

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.

**Directions**

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adults and children 6 years and over	Take 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° and 25°C (68° - 77°F)

### Inactive ingredients

Lactose monohydrate, microcrystalline cellulose, starch (corn), magnesium stearate, hypromellose, polydextrose, polyethylene glycol and titanium dioxide.

### Questions or comments?

**1-800-645-2158**

\*This product is not manufactured or distributed by McNeil-PPC, Inc., distributor of Zyrtec®. Zyrtec® is a registered trademark of UCB Pharma, S.A.

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Distributed by:

Rugby Laboratories  
 31778 Enterprise Drive  
 Livonia, MI 48150  
[www.rugbylaboratories.com](http://www.rugbylaboratories.com)  
 Re-order No. 255553  
 R-126  
 Rev. 06/14

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### Repackaging Information

Please reference the **How Supplied** section listed above for a description of individual tablets. This drug product has been received by Aphenia Pharma - TN in a manufacturer or distributor packaged configuration and repackaged in full compliance with all applicable cGMP regulations. The package configurations available from Aphenia are listed below:

Count	10 mg
90	71610-093-60

Store between 20°-25°C (68°-77°F). See USP Controlled Room Temperature. Dispense in a tight light-resistant container as defined by USP. Keep this and all drugs out of the reach of children.

Repackaged by:



Cookeville, TN 38506

20180717JH

**PRINCIPAL DISPLAY PANEL - 10 mg**

NDC 71610-093 - Cetirizine HCl 10 mg - Rx Only



**CETIRIZINE HYDROCHLORIDE**

cetirizine hydrochloride tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:716 10-093(NDC:0536-1041)
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	10 mg

**Inactive Ingredients**

Ingredient Name	Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
STARCH, CORN (UNII: O8232NY3SJ)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	

<b>POLYDEXTROSE</b> (UNII: VH2XOU12IE)	
<b>POLYETHYLENE GLYCOL 3350</b> (UNII: G2M7P15E5P)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

### Product Characteristics

<b>Color</b>	WHITE	<b>Score</b>	no score
<b>Shape</b>	RECTANGLE (pillow-shaped)	<b>Size</b>	9 mm
<b>Flavor</b>		<b>Imprint Code</b>	10 MG;APO
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71610-093-60	90 in 1 BOTTLE; Type 0: Not a Combination Product	06/28/2018	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078317	12/27/2007	

**Labeler** - Aphena Pharma Solutions - Tennessee, LLC (128385585)

### Establishment

Name	Address	ID/FEI	Business Operations
Aphena Pharma Solutions - Tennessee, LLC		128385585	REPACK(71610-093)

Revised: 7/2018

Aphena Pharma Solutions - Tennessee, LLC