

# CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet

Valu Merchandisers Company

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**Cetirizine Hydrochloride**

***Drug Facts***

**Active ingredient (in each tablet)**

Cetirizine HCl, USP 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 | one 10 mg tablet once daily; do not  
take more than one 10 mg tablet in 24

adults and children 6 years and over	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

- **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.**
- store between 20° to 25° C (68° to 77° F)

### Inactive ingredients

corn starch, hypromellose, lactose monohydrate, magnesium stearate, polyethylene glycol, povidone, talc, titanium dioxide

### Questions?

call **1-800-406-7984**

**PROUDLY DISTRIBUTED BY:  
VALU MERCHANDISERS, CO.  
5000 KANSAS AVE  
KANSAS CITY, KS 66106**

### PRINCIPAL DISPLAY PANEL - 14 Tablet Blister Pack Carton

COMPARE TO THE ACTIVE INGREDIENT IN ZYRTEC®\*

Best  
Choice®

ORIGINAL PRESCRIPTION STRENGTH  
ALLERGY RELIEF

Cetirizine HCl Tablets, USP 10 mg

ANTIHISTAMINE

24 Hour Allergy Relief

Indoor & Outdoor Allergies

Relief of:

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

14 TABLETS 10 mg Each

R0814

All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Zyrtec®.

Keep the carton. It contains important information. See end panel for expiration date.

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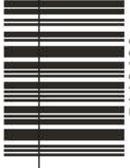
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Best Choice®

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ANTIHISTAMINE

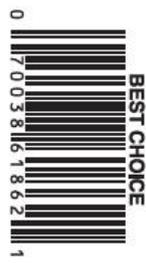
Expiration Date:

Batch No.

Non Varnish Area

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VALU MERCHANDISERS CO.  
5000 KANSAS AVE  
KANSAS CITY, KS 66106



BEST CHOICE

0 7003816 1862 1



## CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63941-939
<b>Route of Administration</b>	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
STARCH, CORN (UNII: O8232NY3SJ)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

### Product Characteristics

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	RECTANGLE (Rounded Off)	<b>Size</b>	9mm
<b>Flavor</b>		<b>Imprint Code</b>	R152
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63941-939-54	14 in 1 BLISTER PACK; Type 0: Not a Combination Product	12/27/2007	
2	NDC:63941-939-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2007	
3	NDC:63941-939-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2007	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077498	12/27/2007	

**Labeler** - Valu Merchandisers Company (868703513)

**Registrant** - Ranbaxy Pharmaceuticals Inc. (937890044)

**Establishment**

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(63941-939)

Revised: 8/2018

Valu Merchandisers Company