CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet
Chain Drug Consortium, LLC

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 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

- **TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL IS BROKEN OR MISSING FROM BOTTLE.** ***(for bottle cartons/stand-alone labels only)***
- **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.** ***(for blister cartons only)***
- 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**

Keep the carton. It contains important information.

See end panel for expiration date.

Distributed By:
Pharmacy Value Alliance, LLC
407 East Lancaster Avenue,
Wayne, PA 19087

PRINCIPAL DISPLAY PANEL - 10 mg Tablet Bottle Carton

COMPARE TO THE ACTIVE
INGREDIENT IN ZYRTEC®*

Original Prescription Strength

Premier
Value®

24 Hour Allergy Relief

Cetirizine HCl Tablets, USP 10 mg

Antihistamine

Indoor & Outdoor Allergies

Relief of:

- Sneezing
- Runny nose
- Itchy, watery eyes
- Itchy throat or nose

90
Tablets
10 mg Each

INDEPENDENTLY TESTED
PV
SATISFACTION GUARANTEED


Original Prescription Strength
Premier Value
24 Hour Allergy Relief
Cetirizine HCl Tablets, USP 10 mg
Antihistamine
Indoor & Outdoor Allergies
90 Tablets 10 mg Each
COMPARE TO THE ACTIVE INGREDIENT IN ZYRTEC®*
Relief of:
✓ Sneezing
✓ Runny nose
✓ Itchy, watery eyes
✓ Itchy throat or nose
Distributed By:
Pharmacy Value Alliance, LLC
407 East Lancaster Avenue,
Wayne, PA 19087
R0817
Expiration Date:
Batch No.
If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.
Non Varnish Area
Drug Facts
Active Ingredient (in each tablet)
Purpose
Cetirizine HCl, USP 10 mg
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.
Keep the carton. It contains important information. See end panel for expiration date.
*All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Zyrtec®.
Drug Facts (continued)
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
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TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL IS BROKEN OR MISSING FROM BOTTLE.
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
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517421

CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-939
Route of Administration	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)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	

Product Characteristics			
Color	white	**Score**	no score
Shape	RECTANGLE (rounded-off)	**Size**	9mm
Flavor		**Imprint Code**	R152
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:68016-939-54	14 in 1 BLISTER PACK; Type 0: Not a Combination Product	12/27/2007	
2	NDC:68016-939-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2007	
3	NDC:68016-939-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2007	
4	NDC:68016-939-90	90 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 - Chain Drug Consortium, LLC (101668460)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	MANUFACTURE(68016-939)

Revised: 8/2018 Chain Drug Consortium, LLC