CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet
Army & Air Force Exchange Service

Drug Facts

ACTIVE INGREDIENT (IN EACH CAPLET)

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 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**

PRINCIPAL DISPLAY PANEL

exchange select

NDC 55301-939-56

†Compare to the active ingredient of Zyrtec®

Original Prescription Strength

ALLERGY RELIEF

CETIRIZINE HCl TABLETS, 10 mg

Antihistamine

Allergy

Indoor & Outdoor Allergies

24 HOUR RELIEF OF

Sneezing

Runny Nose

Itchy, Watery Eyes

Itchy Throat or Nose

5 TABLETS

10 mg EACH

Manufactured For: Your Military Exchanges

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

†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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Non Varnish Area

Batch No.

Expiration Date:

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ALLERGY RELIEF

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exchange select

†Compare to the active ingredient of Zyrtec®

NDC 55301-939-56

6 14299 40367 3

"SATISFACTION GUARANTEED OR YOUR MONEY BACK"

Manufactured For: Your Military Exchanges
By: Ohm Laboratories Inc.
New Brunswick, NJ 08901

5087914

5087914

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:

Drug Facts (continued)

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Inactive ingredients

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CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55301-939
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)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ989GH94E)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

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:55301-939-56	5 in 1 BLISTER PACK		
2	NDC:55301-939-30	30 in 1 BOTTLE		

Marketing Information

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

Labeler - Army & Air Force Exchange Service (001695568)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(55301-939)

Revised: 10/2012 Army & Air Force Exchange Service