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 HCI TABLETS, 10 mg

Antihis tamine

Allergy

Indoor & Outdoor Allergies

24 HOUR RELIEF OF

Sneezing

Runny Nose

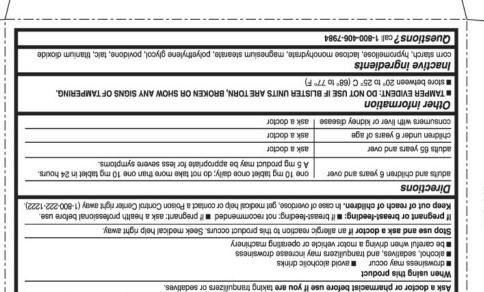
Itchy, Watery Eyes

Itchy Throat or Nose

5 TABLETS

10 mg EACH

Manufactured For: Your Military Exchanges



Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose. Do not nee if you have ever had an allergic reaction to this product or any of its ingredients or to an antihistamine containing

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Cetirizine HCI, USP 10 mg..... Antihistamine Purpose Drug Facts

> exchange select

NDC 55301-939-56

Expiration Date

Non Varnish Area

Warnings

■ Lnuuλ uose

[†]Compare to the active ingredient of Zyrtec®

Drug Facts (continued)

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■ itching of the nose or throat

Original Prescription Strength

ALLERGY RELIEF

CETIRIZINE HCI 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

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

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

"SATISFACTION GUARANTEED OR YOUR MONEY BACK

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



CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:55301-939

Route of Administration ORAL

Active Ingredient/Active Moiety

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

Inactive Ingredients			
Ingredient Name	Strength		
STARCH, CORN (UNII: O8232NY3SJ)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
LACTO SE MO NO HYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)			
PO VIDO NE (UNII: FZ989 GH94E)			
TALC (UNII: 7SEV7J4R1U)			
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)			

Product Characteristics			
Color	white	Score	no score
Shape	RECTANGLE (Rounded-Off)	Size	9 mm
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