

CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet
Medline Industries, Inc.

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

- store between 20° to 25° C (68° to 77° F)
- **TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL IS BROKEN OR MISSING FROM BOTTLE.**

INACTIVE INGREDIENTS

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

QUESTIONS?

Call **1-800-MEDLINE (633-5463)**, Monday - Friday, 9AM - 5PM CST

PRINCIPAL DISPLAY PANEL

MEDLINE

NDC 53329-652-38

†Compare to the active ingredient in Zyrtec®

Original Prescription Strength

CETIRIZINE HCl TABLETS, 10 mg

ANTIHISTAMINE

24 HOUR ALLERGY

Indoor & Outdoor Allergies

Relief of:

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

10 mg

90 TABLETS - 10 mg EACH

Distributed by: Medline Industries, Inc.

5099013/1012



CETIRIZINE HCl TABLETS, 10 mg

ANTI-HISTAMINE
24 HOUR ALLERGY

Indoor &
Outdoor
Allergies



NDC 53329-652-38

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90 TABLETS - 10 mg EACH



8 84389 82339 5

Expiration Date:

Lot No.:

Non Varnish Area

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

ANTI-HISTAMINE
24 HOUR ALLERGY
CETIRIZINE HCl TABLETS, 10 mg



Indoor &
Outdoor
Allergies

Re-order # 07C6817N

1012

Distributed by: Medline Industries, Inc.
One Medline Place, Mundelein, IL 60060

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

Drug Facts (continued)

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

Questions? call 1-800-MEDLINE (633-6463), Monday - Friday, 9AM - 5PM CST

Drug Facts

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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

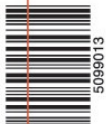
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Ask a doctor or pharmacist before use if you are taking tranquilizers or sedatives.

When using this product
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CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53329-652
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)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
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:53329-652-38	90 in 1 BOTTLE		

Marketing Information

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

Labeler - Medline Industries, Inc. (025460908)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(53329-652)

Revised: 3/2013

Medline Industries, Inc.