

CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet
Select Brand

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 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

PRINCIPAL DISPLAY PANEL

select brand®

NDC 15127-909-30

Original Prescription Strength

Allergy

Cetirizine HCl Tablets, 10 mg

Antihistamine

Indoor & Outdoor Allergies

†Compare to the active ingredient of Zyrtec®

24 Hour Relief of

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

30 Tablets

10 mg EACH

Distributed by: SELECT BRAND® DISTRIBUTORS

5087905/0811



Original Prescription Strength
Allergy
Cetirizine HCl Tablets, 10 mg
Antihistamine
 Indoor & Outdoor Allergies

30 Tablets
10 mg EACH

select brand
 the lower price name brand

NDC 15127-909-30

Original Prescription Strength
Allergy
Cetirizine HCl Tablets, 10 mg
Antihistamine
 Indoor & Outdoor Allergies

†Compare to the active ingredient of Zyrtec®

24 Hour • Sneezing • Runny Nose
Relief of • Itchy, Watery Eyes • Itchy Throat or Nose



30 Tablets
10 mg EACH

Original Prescription Strength
Allergy
Cetirizine HCl Tablets, 10 mg
Antihistamine
 Indoor & Outdoor Allergies

30 Tablets
10 mg EACH



Expiration Date:

Non Varnish Area



Distributed by:
SELECT BRAND® DISTRIBUTORS
 Pine Bluff, AR 71603 USA
 AC (870)-535-3635
 Item # 1008556

Batch No.

0811

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.

Drug Facts (continued)
Inactive ingredients
 corn starch, hypromellose, lactose monohydrate, magnesium stearate, polyethylene glycol, polydioxane, talc, titanium dioxide

Questions? call 1-800-406-7984

Drug Facts
Active ingredient (in each tablet)
 Cetirizine HCl, USP 10 mg.....Antihistamine
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 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
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 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
 ■ avoid alcoholic drinks
 ■ drowsiness may occur
 ■ 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).
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CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:15127-909
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:15127-909-14	14 in 1 BLISTER PACK		
2	NDC:15127-909-30	30 in 1 BOTTLE		
3	NDC:15127-909-90	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 - Select Brand (043562370)**Registrant** - Ranbaxy Pharmaceuticals Inc. (937890044)**Establishment**

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(15127-909)

Revised: 9/2012

Select Brand