

CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet
Publix Super Markets 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

- **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

PRINCIPAL DISPLAY PANEL

Publix

INDOOR & OUTDOOR

allergy relief

CETIRIZINE HYDROCHLORIDE TABLETS, 10 mg

ANTI-HISTAMINE

24 hour relief of:

- sneezing
- itchy, watery eyes
- runny nose
- itchy throat or nose

Original Prescription Strength

60 TABLETS 10 mg EACH

***Compare to Active Ingredient in Zyrtec®**

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5087913/R0711

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INDOOR & OUTDOOR

allergyrelief

CETIRIZINE HYDROCHLORIDE TABLETS, 10 mg

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*Compare to Active
Ingredient in Zyrtec®



Expiration Date:

DISTRIBUTED BY PUBLIX SUPER MARKETS, INC.
3300 PUBLIX CORPORATE PARKWAY
LAKELAND, FL 33811 1-888-267-3037
publix.com

PUBLIX GUARANTEE:
COMPLETE SATISFACTION OR YOUR MONEY BACK

Batch No.

Non Varnish Area

R0711

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

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

Drug Facts (continued)

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

Other Information

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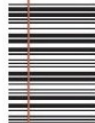
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Uses

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Active ingredient (in each tablet)

Cetirizine HCl, USP 10 mg.....Antihistamine



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CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56062-939
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:56062-939-54	14 in 1 BLISTER PACK		
2	NDC:56062-939-30	30 in 1 BOTTLE		
3	NDC:56062-939-60	60 in 1 BOTTLE		

Marketing Information

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

Labeler - Publix Super Markets Inc (006922009)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(56062-939)

Revised: 12/2012

Publix Super Markets Inc