

**ALLERGY RELIEF- cetirizine hcl capsule**  
**Publix Supermarkets, Inc.**

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**Cetirizine Hydrochloride Tablets**

**Active Ingredient (in each capsule)**

Cetirizine HCl 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

**Warning**

**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
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic drinks
- 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 (1-800-222-1222) right away.

**Directions**

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Adults and children 6 years and over	One 10 mg capsule once daily; do not take more than one 10 mg capsule 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

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**Other Information**

- store at 68° to 77°F (20° to 25°C)
- avoid high humidity and excessive heat above 40° C (104° F)
- protect from light

**Inactive Ingredients**

Butylated hydroxytolene, gelatin, glycerin, mannitol, pharmaceutical ink, polyethylene glycol 400, purified water, sodium hydroxide, sorbitan, sorbitol

**Questions or comments?****Principal Display Panel**

ORIGINAL PRESCRIPTION STRENGTH

**Allergy Relief**

CETIRIZINE HCl CAPSULES 10 mg

ANTIHISTAMINE

24 Hour Relief of:

- Sneezing
- Runny nose
- Itchy, watery eyes
- Itchy throat or nose
- Indoor & outdoor allergies

SOFTGELS\*\*

\*\*LIQUID-FILLED CAPSULES

\*Compare to the active ingredient in Zyrtec®

\*This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Zyrtec®.

**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.**

**KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.**

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1-888-267-3037 [publix.com](http://publix.com)

**Product Label**

**Drug Facts (continued)**

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NDC 56062-741-25

ORIGINAL PRESCRIPTION STRENGTH

**allergyrelief**

**CETIRIZINE HCl CAPSULES 10 mg**

ANTIHISTAMINE

**24-hour relief of:**

- Sneezing
- Runny nose
- Itchy, watery eyes
- Itchy throat or nose
- Indoor & outdoor allergies



ACTUAL SIZE  
\*Compare to the active ingredient in Zyrtec®

**25** SOFTGELS\*\*  
\*\*LIQUID-FILLED CAPSULES



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\*\*LIQUID-FILLED CAPSULES

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Lot No.:  
Exp. Date:

PUBLIX Allergy Relief

## ALLERGY RELIEF

cetirizine hcl capsule

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:56062-741
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNION CARBIDE/COA) (CETIRIZINE	CETIRIZINE	

CETIRIZINE HYDROCHLORIDE (UNII: 640047K10A) (CETIRIZINE - UNII:YO7261ME24)		CETIRIZINE HYDROCHLORIDE	10 mg	
Inactive Ingredients				
Ingredient Name			Strength	
GELATIN (UNII: 2G86QN327L)				
GLYCERIN (UNII: PDC6A3C0OX)				
MANNITOL (UNII: 3OWL53L36A)				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)				
WATER (UNII: 059QF0KO0R)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
SORBITAN (UNII: 6O92ICV9RU)				
SORBITOL (UNII: 506T60A25R)				
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)				
Product Characteristics				
Color	yellow	Score	no score	
Shape	OVAL	Size	13mm	
Flavor		Imprint Code	10	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56062-741-25	1 in 1 BOX	02/28/2019	02/28/2025
1		25 in 1 BOTTLE; Type 0: Not a Combination Product		
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA207235		02/28/2019	02/28/2025

**Labeler** - Publix Supermarkets, Inc. (006922009)