

ALL DAY ALLERGY RELIEF- cetirizine hcl capsule
P & L Development, LLC

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

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

Other Information

- store at 20°- 25° C (68° to 77° F)
- 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?

Call 1-877-753-3935 Monday-Friday 9AM-5PM EST

Principal Display Panel

all-day allergy

cetirizine HCl capsules 10 mg

Antihistamine

sofgels**

(**liquid-filled capsules)

indoor & outdoor allergies

*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.

Distributed by: **PL Developments**

200 Hicks Street, Westbury, NY 11590

Product Label

Exp. Date: _____
 Lot No.: _____
 F0007819
 PLD-B728A
 3 59726 21127 7
 *This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Zyrtec®.

Drug Facts (continued)

- 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.

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WELLNESS BASICS All Day Allergy Relief

ALL DAY ALLERGY RELIEF

cetirizine hcl capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59726-063
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		
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 (Clear)	Score	no score	
Shape	OVAL	Size	13mm	
Flavor		Imprint Code	C10	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59726-063-25	1 in 1 BOX	05/28/2021	
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	ANDA213105	05/28/2021		

Labeler - P & L Development, LLC (800014821)

Revised: 12/2022

P & L Development, LLC