

CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet
LIDL US, LLC

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 BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.**
- 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 or Comments?

1-877-753-3935

DISTRIBUTED BY
Lidl US, LLC
3500 S. Clark Street
Arlington, VA 22202

PRINCIPAL DISPLAY PANEL - 10 mg Tablet Blister Pack Carton

NDC 71141-133-32

Original Prescription Strength

†Compare to the active ingredient in Zyrtec®

Actual Size

***Cetirizine
HCl Tablets,
USP 10 mg***

Antihistamine

ALLERGY

24 Hour Relief of:

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

14 tablets
10 mg EACH

**INDOOR &
OUTDOOR ALLERGIES**

GLUE - N.C.



R0917

This product is not affiliated with the makers/owners of Zyrtec.
All trademarks are property of their respective owners.

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

Drug Facts

Active ingredient (each tablet): Cetirizine HCl, USP 10 mg

Purpose: Antihistamine

Uses: Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: sneezing, runny nose, itchy, watery eyes, itchy throat or nose.

Warnings: Use if you are 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 benzquinizone or sedatives.

When using this product, drowsiness may occur. Avoid alcoholic drinks and a bedtime. Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.

Drug Facts (continued)

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. Ask a doctor before use if you are pregnant or breastfeeding. If pregnant or breastfeeding, not recommended. 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).

Other information: Tablets are white to off-white. Tablets are round with "LID" on one side and "10 mg" on the other. Tablets are not to be crushed or chewed.

Inactive ingredients: croscarmellose sodium, hydroxypropyl methylcellulose, polyethylene glycol, polyethylene glycol, polydioxane, talc, titanium dioxide, croscarmellose sodium, hydroxypropyl methylcellulose, polyethylene glycol, polydioxane, talc, titanium dioxide.

Questions or Comments? 1-877-553-3936

Original Prescription Strength

Cetirizine HCl Tablets, USP 10 mg

Antihistamine ALLERGY

INDOOR & OUTDOOR ALLERGIES

24 Hour Relief of:

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

14 tablets
10 mg EACH

NDC 71141-133-32

Original Prescription Strength

Cetirizine HCl Tablets, USP 10 mg

Antihistamine ALLERGY

INDOOR & OUTDOOR ALLERGIES

24 Hour Relief of:

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

14 tablets
10 mg EACH

†Compare to the active ingredient in Zyrtec®

Actual Size

Original Prescription Strength

Cetirizine HCl Tablets, USP 10 mg

Antihistamine ALLERGY

INDOOR & OUTDOOR ALLERGIES

24 Hour Relief of:

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

14 tablets
10 mg EACH

N.C.

5178573

Exhibition title: Batch No.

Non Varnish Area



5178573

N.C.

DISTRIBUTED BY
Lid US, LLC
3500 S. Clark Street
Arlington, VA 22202
QUESTIONS?
1-844-656-5151
www.lid.com



7710-008 800

CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71141-133
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)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (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:71141-133-32	1 in 1 CARTON	09/01/2017	
1		14 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077498	09/01/2017	

Labeler - LIDL US, LLC (079389709)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(71141-133)

Revised: 1/2018

LIDL US, LLC