CETIRIZINE- cetirizine hydrochloride tablet, film coated
VERITYRX, LLC

Cetirizine

Active ingredient (in each tablet)

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

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
- store between 20-25 °C (68-77 °F)
- do not use if printed foil under cap is broken or missing

Inactive ingredients

corn starch, FD&C blue no. 1 aluminum lake, hypromellose, lactose monohydrate, magnesium stearate, polydextrose, polyethylene glycol, povidone, titanium dioxide, triacetin

Questions or comments?

Any adverse reactions contact FDA (888) 463-6332 or www.FDA.gov/medwatch

Principal Display Panel

VERITYRX

NDC: 83939-0003-03 Qty: 100 TABLETS

CETIRIZINE HCL10MG

*See package insert for indications and dosage schedule.

Store between 20° - 25°C (68° - 77°F)
Keep this and all medications out of the reach of children

83939000303

NDC: 83939-0003-03 Qty: 100 TABLETS

CETIRIZINE HCL10MG

(CETIRIZINE HCL Original NDC: 0904-6717-60)

GTIN: 00383939000335

S/N

Exp.

Lot:

RX

Distributed by: VerityRx 20225 NE 16th Place Miami, FL 33179

Call to Reorder: 844-837-4897

Compare to active ingredient in Zyrtec ®

Original Prescription Strength

Cetirizine

Hydrochloride Tablets, 10 mg/Antihistamine

All Day Allergy Relief

INDOOR & OUTDOOR ALLERGIES

24 HOUR RELIEF OF:

Sneezing

Runny Nose

Itchy, Watery Eyes

Itchy Throat or Nose

CETIRIZINE

cetirizine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83939-0003(NDC:0904-6717)
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)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	white	**Score**	no score
Shape	OVAL	**Size**	10mm
Flavor		**Imprint Code**	4H2
Contains			

Packaging

			Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83939-0003-0	50 in 1 BOX, UNIT-DOSE	11/03/2025	
1		1 in 1 POUCH; Type 0: Not a Combination Product		
2	NDC:83939-0003-2	50 in 1 BOX, UNIT-DOSE	11/03/2025	
2		1 in 1 POUCH; Type 0: Not a Combination Product		
3	NDC:83939-0003-3	100 in 1 BOX, UNIT-DOSE	11/03/2025	
3		1 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078336	11/03/2025	

Labeler - VERITYRX, LLC (097807737)

Revised: 10/2025 VERITYRX, LLC