

CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablets tablet, film coated
Proficient Rx LP

Cetirizine Hydrochloride Tablets

Drug Facts

Active ingredient (in each tablet)

Cetirizine HCl, 10mg

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

Ask a doctor before use if you have

Do not use if you have ever had an allergic reaction to this product or any of its ingredients or to an antihistaminecontaining 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

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

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

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

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° to 25°C (68° to 77°F)

Inactive ingredients

hypromellose, lactose monohydrate, magnesium stearate, polyethylene glycol, povidone, starch, titanium dioxide.

Questions?

call **1-888-375-3784**

Bottle Label



Scan Here



NDC 71205-709-06

Packaged By: Proficient Rx LP
Thousand Oaks, CA 91320



Cetirizine HCl 10mg

#06 Tablets

Each tablet contains: Cetirizine HCl USP, 10 mg
Antihistamine

White, oval shaped, unscored tablets with imprint code "C"

Product ID: QC070906

Dist. By: Dr. Reddy's Laboratories, Inc. Princeton, NJ 08540 Made in India

Store between 20° to 25°C (68° to 77°F)

Keep medication out of the reach of children

Cetirizine HCl 10mg
#06 Tablets SN# MASTER
Lot # 00000 Exp:00/00/00
NDC 71205-709-06

Cetirizine HCl 10mg
#06 Tablets SN# MASTER
Lot # 00000 Exp:00/00/00
NDC 71205-709-06

Cetirizine HCl 10mg
#06 Tablets SN# MASTER
Lot # 00000 Exp:00/00/00
NDC 71205-709-06



GTIN: 00371205709068
SN# MASTER
Exp. 00/00/00
Lot #:00000

CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablets tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71205-709(NDC:43598-811)
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
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	WHITE	Score	no score
Shape	OVAL	Size	7mm
Flavor		Imprint Code	C

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71205-709-06	6 in 1 BOTTLE; Type 0: Not a Combination Product	10/31/2022	
2	NDC:71205-709-10	10 in 1 BOTTLE; Type 0: Not a Combination Product	10/31/2022	
3	NDC:71205-709-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/31/2022	
4	NDC:71205-709-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	10/31/2022	
5	NDC:71205-709-00	100 in 1 BOTTLE; Type 0: Not a Combination Product	07/08/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078343	12/17/2018	

Labeler - Proficient Rx LP (079196022)**Establishment**

Name	Address	ID/FEI	Business Operations
Proficient Rx LP		079196022	REPACK(71205-709) , RELABEL(71205-709)

Revised: 7/2025

Proficient Rx LP