

**CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet
Proficient Rx LP**

Perrigo Cetirizine Hydrochloride Tablets 10 mg Drug Facts

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:

1. runny nose
2. sneezing
3. itchy, watery eyes
4. 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

1. drowsiness may occur
2. avoid alcoholic drinks
3. alcohol, sedatives, and tranquilizers may increase drowsiness
4. 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:

1. if breast-feeding: not recommended
2. 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

1. store between 20 - 25°C (68 - 77°F)
2. 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?

1-800-719-9260

Principal Display Panel

Compare to Zyrtec® active ingredient

Cetirizine Hydrochloride Tablets 10 mg

Antihistamine

Allergy

24 Hour Relief of:

Sneezing

Runny Nose

Itchy, Watery Eyes
 Itchy Throat or Nose
 Original Prescription Strength
 actual size
 30 Tablets
 Indoor & Outdoor Allergies
 NDC 71205-207-30
 Repackaged By:
 Proficient Rx LP
 Thousand Oaks, CA 91320



Scan Here 

NDC 71205-207-30

Packaged By: Proficient Rx LP
 Thousand Oaks, CA 91320

Cetirizine HCl 10mg
#30 Tablets

Each tablet contains: Cetirizine HCl 10 mg
 Antihistamine

White, oval shaped tablet, unscored with imprint code "4H2"

Product ID: QC020730
 Dist. By: Perrigo Allegan, MI 49010
 Store at 20° to 25°C (68° to 77°F)

Keep medication out of the reach of children

Cetirizine HCl 10mg #30 Tablets Lot #:00000 NDC 71205-207-30	SN# MASTER Exp:00/00/00
Cetirizine HCl 10mg #30 Tablets Lot #:00000 NDC 71205-207-30	SN# MASTER Exp:00/00/00
Cetirizine HCl 10mg #30 Tablets Lot #:00000 NDC 71205-207-30	SN#MASTER Exp:00/00/00

GTIN: 00371205207304
 SN# MASTER
 Exp. 00/00/00
 Lot #:00000

CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71205-207(NDC:45802-919)
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)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71205-207-06	6 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2019	
2	NDC:71205-207-10	10 in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2019	
3	NDC:71205-207-15	15 in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2019	
4	NDC:71205-207-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2019	
5	NDC:71205-207-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2019	
6	NDC:71205-207-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078336	12/27/2007	

Labeler - Proficient Rx LP (079196022)

Establishment

Name	Address	ID/FEI	Business Operations
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Revised: 4/2022

Proficient Rx LP