

**CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet, film coated
Preferred Pharmaceuticals Inc.**

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

Indoor & Outdoor Allergies

actual size

Padagis®

Repackaged By: Preferred Pharmaceuticals Inc.

68788-4072

Cetirizine Hydrochloride

Tablet 10mg

Generic for Zyrtec

Each tablet contains Cetirizine HCl 10mg

Pkg Size: Exp Date: #####

Lot#: Batch#:

Ins:

Mfg: Padagis Israel Pharmaceuticals Ltd

Prod#:

Warning

Do not use if you ever had an allergic reaction to this product or any of its ingredients or to an antihistamine containing hydroxyzine. Ask doctor before use if you have liver or kidney disease, or if you are taking sedatives, tranquilizers, or alcohol. While taking this product drowsiness may occur, avoid alcoholic drinks. Store between 20° to 25°C (68° to 77°F). Keep this and all medications out of reach of children. Tablet is oval, white, and imprinted with 4H2.



Directions English

Take _____ tablet(s)
every _____ hours.
Use as directed by your
doctor

CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed.



GTIN

SN #####
EXP #####

Instrucciones Espanol:

Toma _____ tableta(s)
cada _____ horas.
Uso según lo dirigido
por su doctor

Cetirizine Hydrochloride Tablet 10
mg
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Cetirizine Hydrochloride Tablet 10
mg
Qty: Ins:
Lot: Bat:
Prod# (NDC):

Cetirizine Hydrochloride Tablet 10
mg
Qty:
Insurance NDC:
Lot: Bat:

Cetirizine Hydrochloride Tablet 10
mg
Qty: Ins:
Lot: Bat:
Prod# (NDC):

CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-4072(NDC:45802-919)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:Y07261ME24)	CETIRIZINE HYDROCHLORIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: 08232NY3SJ)	
FD&C BLUE NO. 1 ALUMINUM LAKE (UNII: J9EQA3S2JM)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	

Log

Chart

Billing

Patient

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)

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:68788-4072-3	1 in 1 CARTON	01/27/2026	
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:68788-4072-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	01/27/2026	
3	NDC:68788-4072-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	01/27/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA078336	01/27/2026	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-4072)

Revised: 1/2026

Preferred Pharmaceuticals Inc.