

**GOOD SENSE ALL DAY ALLERGY- cetirizine hydrochloride tablet, film coated**  
**Preferred Pharmaceuticals Inc**

-----

**Perrigo All Day Allergy 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:

- 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**

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

**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**

Allergy

Original Prescription Strength

All Day Allergy

Cetirizine Hydrochloride Tablets, 10 mg Antihistamine

Actual Size

24 Hour Relief of:

- Sneezing

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

Indoor & Outdoor Allergies

Compare to active ingredient of Zyrtec®

Repackaged By: Preferred Pharmaceuticals Inc

Cetirizine  
Hydrochloride  
Tablet 10mg

Generic for Zyrtec

Each tablet contains Cetirizine HCl 10mg

Pkg Size:    Exp Date: ####/####/####  
Lot#:            Batch#:

Ins:  
Mfg: L. Perrigo Company  
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 tranquilizers or sedatives. When using this product drowsiness may occur, avoid alcoholic drinks. Store between 20° to 25°C (68° to 77°F). Keep this and all medication out of the reach of children. Tablet is oval, white, and imprinted with 4H2

Directions English

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

GTIN

#####  
SN #####  
EXP ####/####/####

Instrucciones Espanol:

Toma \_\_\_\_\_ tableta(s)  
cada \_\_\_\_\_ horas.  
Usa según lo dirigido por su doctor

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

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:    Ins:  
Insurance NDC:  
Lot:    Bat:

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

Log

Chart

Billing

Patient

GOOD SENSE ALL DAY ALLERGY			
cetirizine hydrochloride tablet, film coated			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-4023(NDC:0113-9458)
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: 3WJQ0SDW1A)			

<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)				
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)				
<b>TRIACETIN</b> (UNII: XHX3C3X673)				
<b>Product Characteristics</b>				
<b>Color</b>	WHITE	<b>Score</b>	no score	
<b>Shape</b>	OVAL	<b>Size</b>	10mm	
<b>Flavor</b>		<b>Imprint Code</b>	4H2	
<b>Contains</b>				
<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:68788-4023-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	09/12/2025	
2	NDC:68788-4023-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	09/12/2025	
3	NDC:68788-4023-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	09/12/2025	
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>		<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ANDA	ANDA078336		09/12/2025	

**Labeler** - Preferred Pharmaceuticals Inc (791119022)

**Registrant** - Preferred Pharmaceuticals Inc (791119022)

Establishment			
Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc		791119022	REPACK(68788-4023)