# **CETIRIZINE HYDROCHLORIDE (ALLERGY) - cetirizine hydrochloride tablet A-S Medication Solutions**

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Cetirizine Hydrochloride Tablets USP 10 mg (ALLERGY RELIEF)

### **Drug Facts**

#### Active ingredient (in each tablet)

Cetirizine hydrochloride USP 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	one 10 mg tablet once daily;
6 years and over	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	ask a doctor
kidney disease	

#### Other information

- store between 20° to 25°C (68° to 77°F)
- TAMPER EVIDENT: DO NOT USE IF SEAL OVER BOTTLE OPENING IS BROKEN OR MISSING

## Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide

#### **Questions?**

call **1-800-206-7821** 

Manufactured for: Northstar Rx LLC

Memphis, TN 38141

Manufactured by: Aurobindo Pharma Limited

Hyderabad-500 090, India

Code: TS/DRUGS/19/1993

Issued: 05/2024

#### **HOW SUPPLIED**

Product: 50090-6170

NDC: 50090-6170-0 30 TABLET in a BOTTLE / 48 in a CASE

#### **Cetirizine Hydrochloride**



## **CETIRIZINE HYDROCHLORIDE (ALLERGY)**

cetirizine hydrochloride tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50090-6170(NDC:16714-799)	
Route of Administration	ORAL			

Active Ingredient/Active Moiety		
Ingredient Name	<b>Basis of Strength</b>	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZ INE HYDROCHLORIDE	10 mg

Inactive Ingredients		
Ingredient Name	Strength	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)		
HYPROMELLOSE 2910 (5 MPA.S) (UNII: R75537T0T4)		

LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	WHITE (White to Off-white)	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	X;36
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:50090- 6170-0	48 in 1 CASE	10/13/2022		
1		30 in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090760	08/05/2015	

# Labeler - A-S Medication Solutions (830016429)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
A-S Medication Solutions		830016429	RELABEL(50090-6170)

Revised: 7/2024 A-S Medication Solutions