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

-----

# **Drug Facts**

#### Active ingredient (in each tablet)

#### For 5 mg:

Cetirizine hydrochloride USP 5 mg

#### For 10 mg:

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**

#### For 5 mg:

adults and children 6 years and over	1 to 2 tablets once daily depending upon severity of symptoms; do not take more than 2 tablets in 24 hours.
adults 65 years and over	1 tablet once a day; do not take more than 1 tablet in 24 hours
children under 6 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

### For 10 mg:

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)
- 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?

Distributed by: **AUROHEALTH LLC** 2572 Brunswick Pike Lawrenceville, NJ 08648

Made in India

Code: TS/DRUGS/19/1993

#### **HOW SUPPLIED**

Product: 50090-6806

NDC: 50090-6806-0 30 TABLET in a BOTTLE / 1 in a CARTON NDC: 50090-6806-1 90 TABLET in a BOTTLE / 1 in a CARTON

### **Cetirizine Hydrochloride**



#### **CETIRIZINE HYDROCHLORIDE (ALLERGY)** cetirizine hydrochloride tablet **Product Information** HUMAN OTC DRUG Item Code (Source) NDC:50090-6806(NDC:58602-445) **Product Type Route of Administration** ORAL **Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE -**CETIRIZ INE** 10 mg UNII:YO7261ME24) **HYDROCHLORIDE Inactive Ingredients Ingredient Name** Strength SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
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- 6806-0	1 in 1 CARTON	11/06/2023	
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:50090- 6806-1	1 in 1 CARTON	11/06/2023	
2		90 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	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-6806), REPACK(50090-6806)

Revised: 1/2024 A-S Medication Solutions