

**CETIRIZINE HYDROCHLORIDE (ALLERGY) - cetirizine hydrochloride tablet**  
**CVS Pharmacy, Inc.**

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**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 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?**

call **1-855-274-4122**

**Distributed by:  
CVS Pharmacy, Inc.**

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Woonsocket, RI 02895  
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CVS.com®  
1-800-SHOP CVS

Made in India

Code: TS/DRUGS/19/1993

**PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 10 mg (30 Tablets  
Container Label)**

♥CVS

Health®

NDC 69842-237-09

Indoor & Outdoor Allergies

Original Prescription Strength Allergy Relief

CETIRIZINE HYDROCHLORIDE TABLETS USP 10 mg Antihistamine

24 Hour 30 TABLETS



\* Lot: XXXXXXXXX  
EXP: MM/YYYY  
Prefix & Variables of Lot, EXP shall be printed online during packing.

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 10 mg (30 Tablets Container Carton Label)

CVS Health®

Compare to the active ingredient in Zyrtec® Tablets\* Indoor & Outdoor Allergies

NDC 69842-237-09

Original Prescription Strength Allergy Relief

CETIRIZINE HYDROCHLORIDE TABLETS USP 10 mg Antihistamine

24 Hour

Relief of:

- Sneezing
- Runny nose
- Itchy, watery eyes
- Itchy throat or nose

Actual Size  
30 TABLETS

Actual Bottle Size  
on Side Panel



## CETIRIZINE HYDROCHLORIDE (ALLERGY)

cetirizine hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69842-237
<b>Route of Administration</b>	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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE 2910 (5 MPA.S) (UNII: R75537T0T4)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q815X)	
MAGNESIUM STEARATE (UNII: 70097M6130)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

### Product Characteristics

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

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69842-237-91	1 in 1 CARTON	07/26/2019	06/10/2023
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:69842-237-09	1 in 1 CARTON	07/26/2019	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:69842-237-17	1 in 1 CARTON	07/26/2019	
3		45 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:69842-237-54	1 in 1 CARTON	07/26/2019	01/11/2021
4		70 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:69842-237-19	1 in 1 CARTON	07/26/2019	
5		90 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:69842-237-23	1 in 1 CARTON	07/26/2019	
6		120 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:69842-237-45	2 in 1 CARTON	07/26/2019	
7		120 in 1 BOTTLE; Type 0: Not a Combination Product		
8	NDC:69842-237-39	365 in 1 BOTTLE; Type 0: Not a Combination Product	07/26/2019	
9	NDC:69842-237-60	1 in 1 CARTON	07/26/2019	01/10/2022
9		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
10	NDC:69842-237-57	1 in 1 CARTON	07/26/2019	01/08/2023
10		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
11	NDC:69842-237-01	1 in 1 CARTON	07/26/2019	
11		14 in 1 BLISTER PACK; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090760	07/26/2019	

**Labeler** - CVS Pharmacy, Inc. (062312574)

**Registrant** - Aurohealth LLC (078728447)

**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Aurobindo Pharma Limited		918917642	ANALYSIS(69842-237) , MANUFACTURE(69842-237)

Revised: 3/2024

CVS Pharmacy, Inc.