#### CETIRIZINE HYDROCHLORIDE (ALLERGY) - cetirizine hydrochloride tablet INNOVUS PHARMACEUTICALS, INC.

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#### Cetirizine Hydrochloride Tablets (Allergy)

**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: Innovus Pharmaceuticals, Inc. Reno, NV 89511 **www.crclehealth.com** 

Made in India

# PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 10 mg (365's Tablet Container Label)

\*Compare to the active ingredient of Zyrtec<sup>®</sup>

#### C•rcle™

NDC 57483-190-14

Allergy Relief Cetirizine Hydrochloride Tablets USP, 10 mg Antihistamine Original Prescription Strength

INDOOR & OUTDOOR ALLERGIES 10 mg Strength Tablets

24

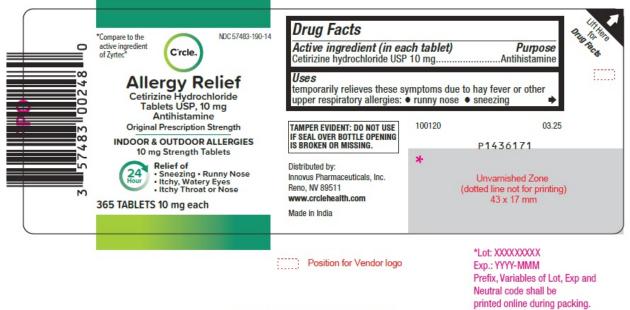
Hour

**Relief of** 

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

365 TABLETS 10 mg each

# Top Layer Printing side



## Top Layer Adhesive side

Drug Facts (continued)	Drug Facts (con	tinued)	
• itchy, watery eyes • itching of the nose or throat	<ul> <li>Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.</li> <li>If pregnant or breast-feeding: <ul> <li>If breast-feeding: not recommended</li> <li>if pregnant: ask a health professional before use.</li> </ul> </li> <li>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]</li> </ul>		
<i>Warnings</i> <b>Do not use</b> if you have ever had an allergic reaction to this product or any of its ingredients or to an antihistamine containing hydroxyzine.			Gluing Area
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.	Directions adults and children	one 10 mg tablet once daily; do	Ū
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	6 years and over	not take more than one 10 mg tablet in 24 hours. A 5 mg product may be appropriate for less severe symptoms.	

## Base Layer

P1436171

Drug Facts (continued)		*This product is not manufactured or distributed by McNeil Consumer Healthcare, Division of McNeil-PPC, Inc.,
adults 65 years and over	ask a doctor	owner of the registered trademark Zyrtec <sup>®</sup> .
children under 6 years of age	ask a doctor	
consumers with liver or kidney disease	ask a doctor	
<ul> <li>store between 20° to 25°C (68° to 77°F</li> </ul>	) Al over bottle opening	
	7	
Questions? call 1-855-274-4122		+ C + 30 / F 0
	adults 65 years and over children under 6 years of age consumers with liver or kidney disease Other information • store between 20° to 25°C (68° to 77°F • TAMPER EVIDENT: DO NOT USE IF SE IS BROKEN OR MISSING. Inactive ingredients colloidal silicon dioxide, croscarmellose so monohydrate, magnesium stearate, micro- polyethylene glycol, titanium dioxide	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       astronom         • 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

# **CETIRIZINE HYDROCHLORIDE (ALLERGY)**

cetirizine hydrochloride tablet

Product Type		HUMAN OTC DRUG Item Code (Source)		NDC:57483-190		
					NDC.5740	55-190
Route of Admin	istration	ORAL				
Active Ingred	ient/Active	Moietv				
Ingredient Name Basis of Stree						Streng
CETIRIZINE HYDR JNII:YO7261ME24)	•	NII: 640047KTOA) (CETIRIZIN	E -	CETIRIZ INE HYDROCHLORID		10 mg
Inactive Ingre	dients					
		Ingredient Name			St	rength
SILICON DIOXIDE	(UNII: ETJ7Z6XB	U4)				
CROSCARMELLOS	E SODIUM (UNI	I: M28OL1HH48)				
		(UNII: R75537T0T4)				
LACTOSE MONOH						
MAGNESIUM STEARATE (UNII: 70097M6I30)						
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)						
POLYETHYLENE G	LYCOL 400 (UN	III: B697894SGQ)				
POLYETHYLENE G TITANIUM DIOXID	E (UNII: 15FIX9V)	III: B697894SGQ)				
POLYETHYLENE G TITANIUM DIOXID Product Chara	E (UNII: 15FIX9V)	III: B697894SGQ) 2JP)	Score		no s	core
POLYETHYLENE G TITANIUM DIOXID <b>Product Char</b> a Color	E (UNII: 15FIX9V2	III: B697894SGQ) 2JP)	Score		no s 8mn	
POLYETHYLENE G TITANIUM DIOXID <b>Product Char</b> a Color Shape	E (UNII: 15FIX9V CUNII: 15FIX9V CONTENTION	III: B697894SGQ) 2JP)		t Code		า
MICROCRYSTALLI POLYETHYLENE G TITANIUM DIOXID <b>Product Char</b> Color Shape Flavor Contains	E (UNII: 15FIX9V CUNII: 15FIX9V CONTENTION	III: B697894SGQ) 2JP)	Size	t Code	8mn	า
POLYETHYLENE G TITANIUM DIOXID <b>Product Chara</b> Color Shape Flavor	E (UNII: 15FIX9V CUNII: 15FIX9V CONTENTION	III: B697894SGQ) 2JP)	Size	t Code	8mn	า
POLYETHYLENE G FITANIUM DIOXID Product Chara Color Shape Flavor Contains Packaging	E (UNII: 15FIX9V)	III: B697894SGQ) 2JP)	Size	t Code eting Start Date	8mn X;36 Market	1
POLYETHYLENE G TITANIUM DIOXID Product Chara Color Shape Flavor Contains Packaging # Item Code	E (UNII: 15FIX9V)	III: B697894SGQ) 2JP) to Off-white)	Size Imprint Marke	eting Start Date	8mn X;36 Market	ting End
POLYETHYLENE G TITANIUM DIOXID Product Chara Color Shape Flavor Contains Packaging # Item Code 1 NDC:57483-190-	E (UNII: 15FIX9V) ACTERISTICS WHITE (White ROUND Pac 365 in 1 BOTTI	III: B697894SGQ) 2JP) to Off-white) <b>kage Description</b>	Size Imprin Marke	eting Start Date	8mn X;36 Market	ting End
POLYETHYLENE G FITANIUM DIOXID Product Chara Color Shape Flavor Contains Packaging # Item Code 1 NDC:57483-190- 14	E (UNII: 15FIX9V) ACTERISTICS WHITE (White ROUND Pac 365 in 1 BOTTI Product	III: B697894SGQ) 2JP) to Off-white) <b>kage Description</b> LE; Type 0: Not a Combinatio	Size Imprin Marke	eting Start Date	8mn X;36 Market	ting End
POLYETHYLENE G TITANIUM DIOXID Product Chara Color Shape Flavor Contains Packaging # Item Code 1 NDC:57483-190-	E (UNII: 15FIX9V) ACTERISTICS WHITE (White ROUND Pac 365 in 1 BOTTI Product Informati	III: B697894SGQ) 2JP) to Off-white) <b>kage Description</b> LE; Type 0: Not a Combinatio	Marke	eting Start Date	8mn X;36 Market D	ting End

Labeler - INNOVUS PHARMACEUTICALS, INC. (962507187)

Registrant - Aurohealth LLC (078728447)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
APL HEALTHCARE LIMITED		650844777	ANALYSIS(57483-190), MANUFACTURE(57483-190)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
Aurobindo Pharma Limited		918917642	ANALYSIS(57483-190), MANUFACTURE(57483-190)

Revised: 4/2025

INNOVUS PHARMACEUTICALS, INC.