CETIRIZINE HYDROCHLORIDE (ALLERGY)- cetirizine hydrochloride tablet
BluePoint Laboratories

CETIRIZINE HCL Tablets USP

Drug Facts

Active ingredient (in each tablet)

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 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?

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

Manufactured by:

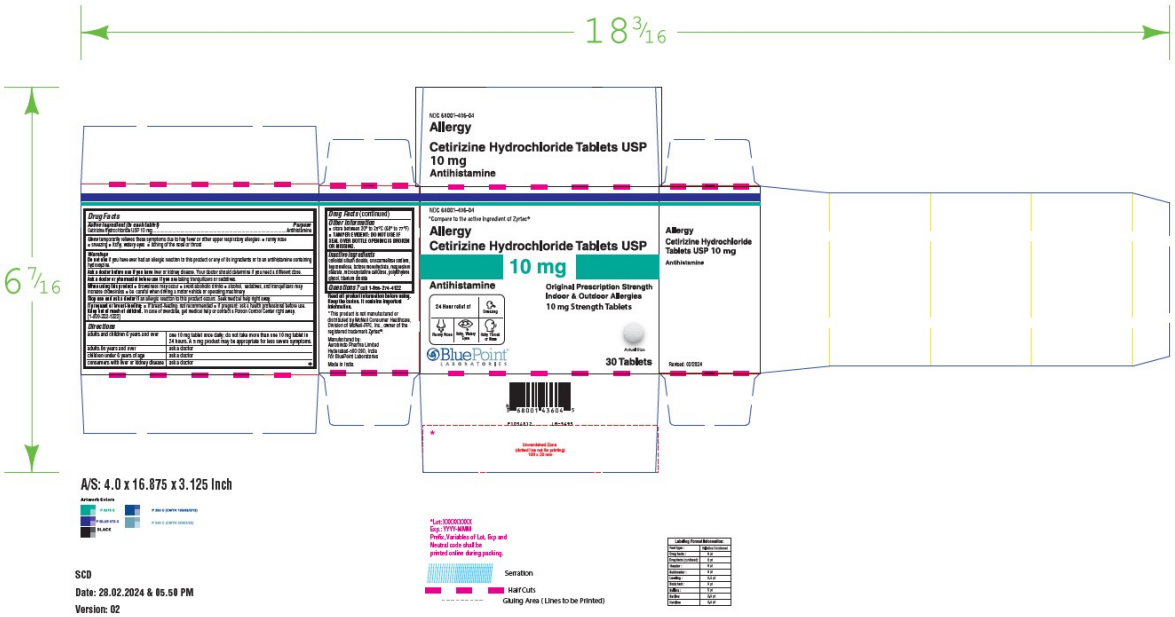
Aurobindo Pharma Limited
Hyderabad-500 090, India
For BluePoint Laboratories

Made in India

Code: TS/DRUGS/19/1993

Issued 05/2020

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 10 mg (30's Tablet Container Carton Label)



NDC 68001-436-04

***Compare to the active**

ingredient of Zyrtec ®

Allergy Relief

Cetirizine Hydrochloride Tablets USP 10 mg

Antihistamine

Original Prescription Strength

Indoor & Outdoor Allergies

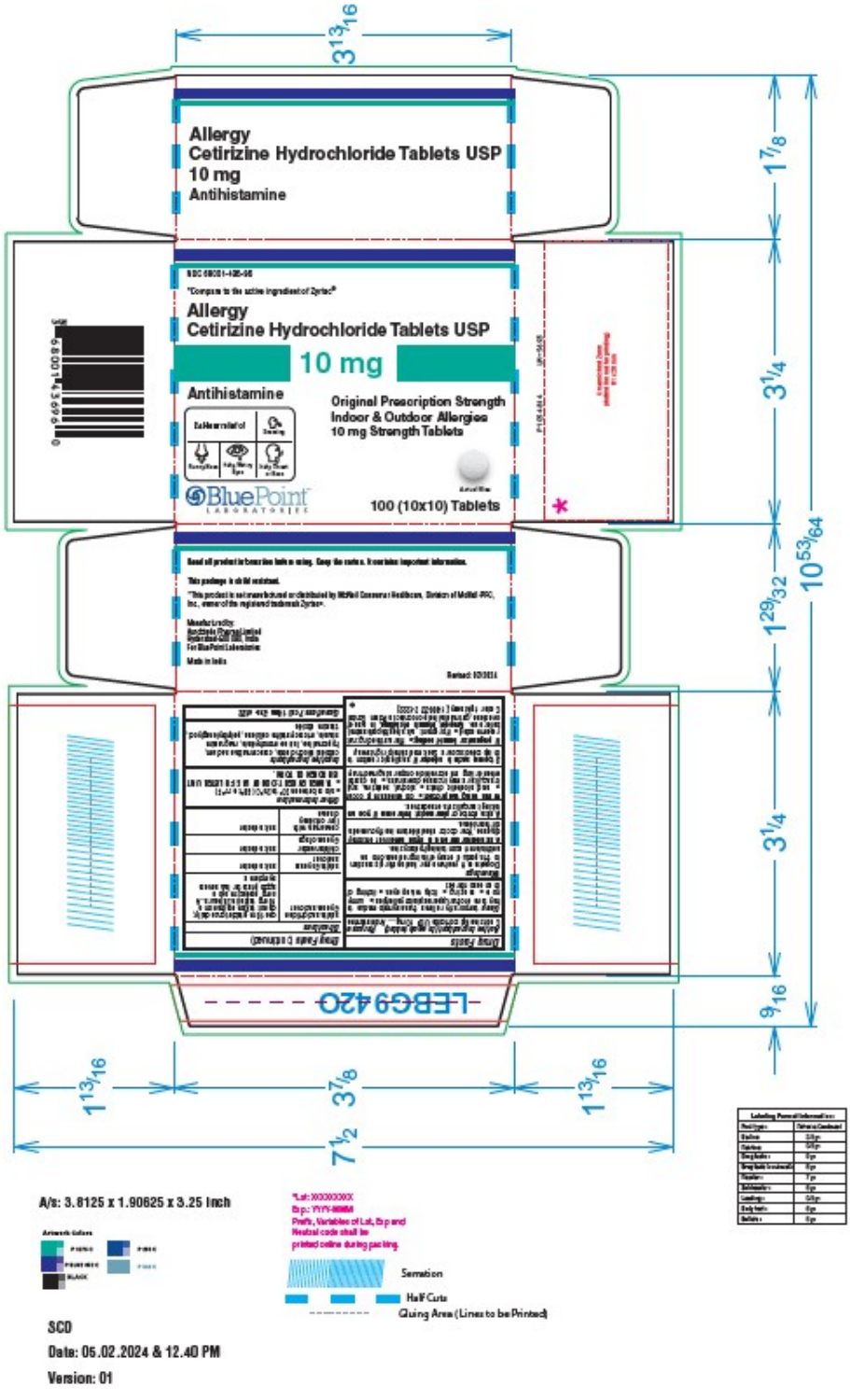
24 Hour Relief of :

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

30 Tablets

10 mg each

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 10 mg (10 x 10 Blister Carton Label)



NDC68001-436-96

***Compare to the active**

ingredient of Zyrtec ®

Allergy Relief

Cetirizine Hydrochloride Tablets USP 10 mg

Antihistamine

Original Prescription Strength

Indoor & Outdoor Allergies

24 Hour Relief of :
- **Sneezing**
- **Runny Nose**
- **Itchy, Watery Eyes**
- **Itchy Throat or Nose**

100 (10x10) Tablets

10 mg each

CETIRIZINE HYDROCHLORIDE (ALLERGY)

cetirizine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68001-436
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**
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			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68001-436-96	10 in 1 CARTON	06/19/2020	
1	NDC:68001-436-16	10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:68001-436-04	1 in 1 CARTON	06/19/2020	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:68001-436-97	1 in 1 CARTON	06/19/2020	
3		300 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	06/19/2020	

Labeler - BluePoint Laboratories (985523874)

Establishment

Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		918917642	analysis(68001-436) , manufacture(68001-436)

Revised: 10/2025 BluePoint Laboratories