CETIRIZINE HYDROCHLORIDE - cetirizine hydrochloride tablet, orally disintegrating WALGREEN CO.

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

Tablet melts in mouth. Can be taken with or without water.

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). Avoid high humidity.
- do not use if carton or blister unit is opened or broken
- see side panel for lot number and expiration date

Inactive ingredients

betadex, citric acid anhydrous, colloidal silicon dioxide, crospovidone, dl-alphatocopherol, hydroxypropyl cellulose, magnesium stearate, maize maltodextrin, mannitol, microcrystalline cellulose, natural flavourings, sodium bicarbonate, sodium starch glycolate and sucralose.

Questions or comments?

call **1-855-274-4122** (Monday - Friday 8:30 AM to 5:00 PM EST)

DISTRIBUTED BY: WALGREEN CO.

200 WILMOT RD., DEERFIELD, IL 60015

MADE IN INDIA

Code: AP/DRUGS/04/2016

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL -10 mg (24 Orally Disintegrating Tablets) Blister Carton

Walgreens

Compare to the active ingredient of Zyrtec[®] Allergy^{††} **NDC 0363-0471-76**

ORIGINAL PRESCRIPTION STRENGTH
Allergy Relief
24 HOUR ALLERGY
CETIRIZINE HYDROCHLORIDE ORALLY DISINTEGRATING TABLETS, USP 10
mg/ANTIHISTAMINE

24 Hour Dissolve Tabs Indoor & Outdoor Allergies

- Relief of sneezing; runny nose;
 itchy, watery eyes & itchy throat or nose
- Melts in your mouth

24 ACTUAL SIZE Orange HOUR flavor

24 ORALLY
DISINTEGRATING
TABLETS
10 mg each





24 HOUR ALLE<mark>r</mark>gy Cet rzine hydrogi orde opally disintegrating tabletş lib 10 mg /ant h'yan ne



N DC0 383+04 1.1-1.8

ORIGINAL PRESCRIPTION STRENGTH

24 HOOR ALLERGY CHIRIMEHORICKHORIDG ORALLYOS IMEGRATING TARLES, USP 10 mg /A MIHISTAMINE



¥ 5

LM-4618

P1044680

Drug Facts

Active ingredient (in each tablet) Purpose Cetirizine hydrochloride USP 10 mg

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

Do not use if you have ever had an allergic reaction to this product or any of its ingredients or to an antihistamine

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 tranquitizers or sedatives.

When using this product

drowstress may occur = avoid alcoholic drinks

alcohol, sedatives, and tranquitizers 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

Drug Facts (continued)

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

Tablet melts in mouth. Can be taken with or without water

one 10 mg tablet once daily; do not take more than one 10 mg tablet in 24 hours. A 5 mg product may be adults and children 6 years and over appropriate for less severe symptoms

adults 6δ years and over ask a doctor children under ask a doctor 6 years of age ask a doctor or kidney disease

Other information

store between 20° to 25°C (68° to 77°F). Avoid high humidity.
 do not use if carton or blister unit is opened or broken
 see side panel for lot number and expiration date.

Inactive ingredients betadex, citric acid anhydrous, colloidal materier injectionist occasio, cinci acia arriyatosa, conocia sistoni diccide, crosposidone, di-alpha-teopheral, hydroxyropyl cellulose, magnesium stearate, maize maltodextrin, mannitol, microcrystalline cellulose, natural librourings, sodium bicarbonate, sodium starch glycolate and sucralose. W0000-0000-0 TEM 7 10955

W30RG0821-F

DISTRIBUTED SY WALGREN CO.
ZON WILL WOOD RO, DEEPFIELD, IL 60015
100% SATISFACTION GUARANTEED
WAIGMENT.COM
62021 WAIGMEN CO. MADE IN INDIA

Code: APIDRUGS04/2010

Drug Facts (continued)

Questions or comments? call 1-855-274-4122 (Monday – Friday 8:30 AM to 5:00 PM EST)

*Our pharmacists recommend the Walgreens brand. We invite you to compare to national brands.
This product is not manufactured or distributed by Johnson & Johnson Inc., MCNeil Consumer Healthcare Division, owner of the registered trademark Zyrtec Allergy.



CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet, orally disintegrating

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0363-0471

Route of Administration ORAL

Active Ingredient/Active Moiety

Active ingredient/Active Plotety			
Ingredient Name	Basis of Strength	Strength	
CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZ INE HYDROCHLORIDE	10 mg	

Inactive Ingredients		
Ingredient Name	Strength	
BETADEX (UNII: JV039JZZ3A)		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
CROSPOVIDONE (35 .MU.M) (UNII: 40UAA97IT9)		
.ALPHATOCOPHEROL, DL- (UNII: 7QWA1RIO01)		
HYDROXYPROPYL CELLULOSE (110000 WAMW) (UNII: 5Y0974F5PW)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MALTODEXTRIN (UNII: 7CVR7L4A2D)		
MANNITOL (UNII: 3OWL53L36A)		
MICROCRYSTALLINE CELLULOSE 101 (UNII: 7T9FYH5QMK)		
MICROCRYSTALLINE CELLULOSE 102 (UNII: PNR0YF693Y)		
SODIUM BICARBONATE (UNII: 8MDF5V39QO)		
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)		
SUCRALOSE (UNII: 96K6UQ3ZD4)		

Product Characteristics			
Color	WHITE	Score	no score
Shape	ROUND	Size	10mm
Flavor	ORANGE	Imprint Code	CE;10
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0471- 76	4 in 1 CARTON	09/11/2020	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph	Marketing Start	Marketing End

ANDA	ANDA213557	09/11/2020

Labeler - WALGREEN CO. (008965063)

Registrant - Aurohealth LLC (078728447)

Establishment			
Name	Address	ID/FEI	Business Operations
APL HEALTHCARE LIMITED		650918514	ANALYSIS(0363-0471), MANUFACTURE(0363-0471)

Revised: 9/2021 WALGREEN CO.