CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet ST. MARY'S MEDICAL PARK PHARMACY

Cetirizine Hydrochloride Tablets USP, 10 mg, Allergy

ACTIVE INGREDIENTS

Active Ingredients (in each tablet)

Purpose

Cetirizine HCI USP 10 mg......Antihistimine

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 DOCTOR

Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.

ASK DOCTOR/PHARMACIST

Ask a doctor or pharmacist before use if you are taking tranquilizers or sedatives.

WHEN USING THIS PRODUCT

- drowsines may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinary.

STOP USE

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 Poison Control Center right away.

DIRECTIONS

Adults and	one 10 mg tablet once daily, do not take more than one 10
children 6	mg tablet in 24 hours. A 5 mg product may be appropriate
years and over	for less severe symptoms.
Adults 65 years	Ask a doctor
and over	ASK & GOCTOL
Children under	Ask a doctor
6 years of age	ASK & GUCLUI
Consumers	
with liver or	Ask a doctor
kidney disease	

OTHER INFORMATION

store at 20° to 25°C (68° to 77°F)

[See USP Controlled Room Temperature]

INACTIVE INGREDIENTS

Hypromellose, lactose, magnesium stearate, maize starch, polyethylene glycol, povidone, titanium dioxide.

QUESTIONS?

Call 1-844-874-7464

Manufactured by:

Unique Pharmaceutical Labs.

(A Div. of J. B. Chemicals & Pharmaceuticals Ltd.), Mumbai 400 030, India

Distributed by:

Rising Pharma holdings, Inc. East Brunswick, NJ 08816

M. L. G/1430 Jul. 2020

126406

PRINCIPAL DISPLAY PANEL-100'S COUNT



Cetirizine
Hydrochloride
Tablets USP
10mg
QTY: 30
RX# ???????
NDC 60760-886-30
LOT# XXXXXXX
EXP XX-XX

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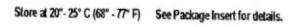
EXP XX-XX

Tablets USP 10mg QTY: 30 RX# ???????? NDC 60760-886-30 LOT# XXXXXXX EXP XX-XX

Cetirizine

Hydrochloride

USE AS DIRECTED





CETIRIZINE HYDROCHLORIDE

ORO VALLEY, AZ 85737

cetirizine hydrochloride tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60760-886(NDC:16571-402)	
Route of Administration	ORAL			

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

Inactive Ingredients		
	Ingredient Name	Strength

HYPROMELLOSES (UNII: 3NXW29V3WO)	
LACTOSE (UNII: J2B2A4N98G)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	
POLYETHYLENE GLYCOL (UNII: 3WJQ0SDW1A)	
POVIDONE (UNII: FZ 989GH94E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics				
Color	white (White)	Score	no score	
Shape	BULLET (Barrel Shaped)	Size	8mm	
Flavor		Imprint Code	CTN;10	
Contains				

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:60760- 886-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/17/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077829	01/17/2023	

Labeler - ST. MARY'S MEDICAL PARK PHARMACY (063050751)

Establishment				
Name	Address	ID/FEI	Business Operations	
ST. MARY'S MEDICAL PARK PHARMACY		063050751	relabel(60760-886), repack(60760-886)	

Revised: 12/2025 ST. MARY'S MEDICAL PARK PHARMACY