

CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet
FAMILY DOLLAR SERVICES, INC

Drug Facts

ACTIVE INGREDIENT (IN EACH TABLET)

Cetirizine HCl, 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

- **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.**
- store between 20° to 25° C (68° to 77° F)

INACTIVE INGREDIENTS

Corn starch, hypromellose, lactose monohydrate, magnesium stearate, polyethylene glycol, povidone, talc, titanium dioxide

QUESTIONS?

Call **1-800-406-7984**

PRINCIPAL DISPLAY PANEL - 10 mg Tablet Blister Pack Carton

COMPARE TO THE ACTIVE INGREDIENT OF ZYRTEC®*

NDC 55319-939-54

FAMILY
wellness™

100% SATISFACTION
OR YOUR MONEY BACK
GUARANTEED

Allergy Relief
Cetirizine HCl Tablets, USP 10 mg/Antihistamine
Indoor & Outdoor Allergies

Relief of:

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

24
HOUR

Original Prescription Strength

14 TABLETS 10 mg EACH

ACTUAL SIZE

OTC
network

Keep the carton. It contains important information. See end panel for expiration date.

Drug Facts (continued)

Active Ingredient
(in each tablet)
Cetirizine HCl, USP 10 mg
Antihistamine

Purpose
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 ■ itchy, watery eyes
 ■ itching of the nose or throat

Warnings
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 When using this product
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 ■ alcohol, sedatives, and tranquilizers may increase drowsiness
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 ■ itchy, watery eyes
 ■ sneezing
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COMPARE TO THE ACTIVE INGREDIENT OF ZYRTEC®

FAMILY wellness™

Allergy Relief

Cetirizine HCl Tablets, USP 10 mg/Antihistamine

COMPARE TO THE ACTIVE INGREDIENT OF ZYRTEC®

NDC 55310-030-54

FAMILY wellness™

100% SATISFACTION OR YOUR MONEY BACK GUARANTEED

Allergy Relief

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Indoor & Outdoor Allergies

Relief of:

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Original Prescription Strength

24 HOUR

ACTUAL SIZE

OTC network

14 TABLETS 10 mg EACH

Cetirizine HCl Tablets, USP 10 mg/Antihistamine

FAMILY wellness™

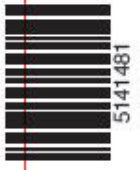
COMPARE TO THE ACTIVE INGREDIENT OF ZYRTEC®

5141481

Expiration Date:

Batch No.

Non Varnish Area



DIS TRIBUTED BY: MIDWOOD BRANDS, LLC
 10611 MONROE RD., MATTHEWS, NC 28105
 NOT 100% SATISFIED?
 Return package and unused product
 within 30 days to any Family Dollar store
 for a refund (with receipt) or exchange.



SKU 0916887

COMPARE TO THE ACTIVE INGREDIENT OF ZYRTEC®

CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55319-939
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
STARCH, CORN (UNII: O8232NY3SJ)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	Score	no score
Shape	RECTANGLE (Rounded Off)	Size	9mm
Flavor		Imprint Code	R152
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55319-939-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2017	
2	NDC:55319-939-54	14 in 1 BLISTER PACK; Type 0: Not a Combination Product	09/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077498	09/01/2017	

Labeler - FAMILY DOLLAR SERVICES, INC (024472631)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(55319-939)

Revised: 1/2018

FAMILY DOLLAR SERVICES, INC