

ALLERGY RELIEF- cetirizine hcl tablet
Strategic Sourcing Services LLC

Drug Facts

Active ingredient (in each tablet)

Cetirizine HCl 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 (1-800-222-1222) right away.

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)
- contains no ingredient made from a gluten-containing grain(wheat, barley or rye)

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide

Questions or comments?

Call **1-833-358-6431** Monday to Friday 9am to 7 pm EST

Principal Display Panel

COMPARE TO ZYRTEC® ACTIVE INGREDIENT*

ORIGINAL PRESCRIPTION STRENGTH

All Day Allergy

Cetirizine HCl 10 mg

ANTIHISTAMINE

Gluten Free

Indoor % Outdoor Allergies

24-Hour symptom relief

- Sneezing
- Runny nose
- Itchy, watery eyes
- Itchy throat or nose

TABLETS

*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division, distributor of Zyrtec®.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPEN OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

Distributed by: McKesson Corp., via Strategic

Sourcing Services LLC. Memphis, TN 38141

www.fosterandthrive.com

Package Labeling

*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division, distributor of Zyrtec®.

Questions or comments?
Call 833-358-6131 Monday to Friday 9:00am to 7:00pm EST

Inactive ingredients colloidal silicon dioxide, croscarmellose sodium, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide

Other information
■ store between 20° to 25°C (68° to 77°F)
■ contains no ingredient made from a gluten-containing grain (wheat, barley or rye)

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.
adults 65 years and over	ask a doctor
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Drug Facts (continued)

Drug Facts
Active ingredient (in each tablet) Cetirizine HCl 10 mg.....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 ■ avoid alcoholic drinks

■ drowsiness may occur ■ 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 pregnant: ask a health professional before use

■ If pregnant: 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 (1-800-222-1222) right away.

Foster & Thrive™
ORIGINAL PRESCRIPTION STRENGTH
All Day Allergy
Cetirizine HCl 10 mg
ANTIHISTAMINE

NDC 70677-1047-1

COMPARE TO ZYRTEC® ACTIVE INGREDIENT*

Foster & Thrive™
ORIGINAL PRESCRIPTION STRENGTH
All Day Allergy
Cetirizine HCl 10 mg
ANTIHISTAMINE

Gluten Free
Indoor & Outdoor Allergies
24-hour symptom relief

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



ACTUAL SIZE

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.
KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

14 TABLETS

Lot No.
Exp. Dt

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Distributed by: McKesson Corp., via Strategic
Sourcing Services LLC, Memphis, TN 38141
Money Back Guarantee
www.fosterandthrive.com

Product of India
PLD-A539A FC008532 Rev 02/23

**FOSTER & THRIVE All Day Allergy Relief****ALLERGY RELIEF**

cetirizine hcl tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-1047
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
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

Product Characteristics

Color	white	Score	2 pieces
Shape	OVAL	Size	9mm
Flavor		Imprint Code	G4
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:70677-1047-1	14 in 1 CARTON	03/01/2023	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA209274	03/01/2023	

Labeler - Strategic Sourcing Services LLC (116956644)

Revised: 5/2024

Strategic Sourcing Services LLC