

ZYRTEC ALLERGY- cetirizine hydrochloride tablet, film coated
Morning Star OTC

Zyrtec[®] Allergy

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 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

- store between 20° to 25°C (68° to 77°F)
- **do not use if foil inner seal imprinted with “Sealed For Your Safety” is broken or missing**
- meets USP Dissolution Test 2

Inactive ingredients

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

Questions?

call **1-800-343-7805**(toll-free) or **215-273-8755**(collect)

Repackaged by:

Morning Star OTC

145 S. Anderson St, Los Angeles,

CA 90033

PRINCIPAL DISPLAY PANEL-1 Tablets

Active ingredient made in Switzerland
Distributed by: JOHNSON & JOHNSON CONSUMER INC.
McNeil Consumer Healthcare Division
Fort Washington, PA 19034 USA
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Pat. www.jjcipts.com
The trade dress of this ZYRTEC® package
is subject to trademark protection.

This product is repackaged by: Morning Star OTC
145 S. Anderson St. Los Angeles, CA 90033
Hours of operation 9 am – 4 pm, 323.354.4838



DO NOT USE IF INNER POUCH OR CARTON IS OPEN OR DAMAGED!

Drug Facts	
Active ingredient (in each caplet)	Purpose
Cetirizine HCl 10mg	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)	

Drug Facts (continued)	
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	
■ do not use if pouch is torn or damaged	■ store between 20 to 25°C (68 to 77°F)
■ meets USP Dissolution Test 2	
Inactive ingredients	
colloidal silicon dioxide, croscarmellose sodium, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide	
Question?	
call 1-800-343-7005 (toll-free) or 215-273-8755 (collect)	

ZYRTEC ALLERGY

cetirizine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53209-5001(NDC:50580-726)
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, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	Score	2 pieces
Shape	RECTANGLE (rounded-off rectangular biconvex tablet)	Size	9mm
Flavor		Imprint Code	ZYRTEC;10;MG
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53209-5001-1	1 in 1 BLISTER PACK; Type 0: Not a Combination Product	06/19/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA019835	06/19/2025	

Labeler - Morning Star OTC (078589357)

Registrant - Morning Star OTC (078589357)

Establishment

Name	Address	ID/FEI	Business Operations
Morning Star OTC		078589357	repack(53209-5001)

Revised: 6/2025

Morning Star OTC