CABINET ALLERGY RELIEF- cetirizine hydrochloride tablet, film coated
Spirit Pharmaceutical LLC

Allergy Relief

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)

Inactive ingredients

colloidal silicon dioxide*, croscarmellose sodium*, hypromellose, lactose, magnesium stearate, maize starch*, microcrystalline cellulose*, polyethylene glycol, povidone*, titanium dioxide

*containes one or more of these ingrdients

Questions or comments?

1-888-333-9792

PRINCIPAL DISPLAY PANEL

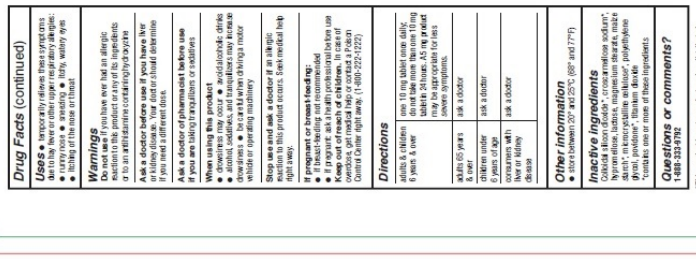
Compare to the active ingredient
in Zyrtec®*

Allergy Relief

Cetirizine HCl 10mg

100 Tablets

*This product is not manufactured or distributed
by McNeil Consumer Healthcare, owner of the
registered trademark Zyrtec®



Drug Facts (continued)

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. The doctor should determine if you should use this product.

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

When using this product ● 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: ask a health professional before use

● 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 & children 6 years & over	one 10 mg tablet once daily; do not take more than one 10 mg tablet in 24 hours; AS mg product may be appropriate for less severe symptoms
adults 65 years & over	ask a doctor
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consumers with liver or kidney disease	ask a doctor

Other information

● store between 20° and 25°C (68° and 77°F)

Inactive ingredients

Colloidal silicon dioxide*, croscarmellose sodium*, hypromellose, lactose monohydrate, magnesium stearate*, microcrystalline cellulose*, polyethylene glycol, povidone*, titanium dioxide

*contains one or more of these ingredients

Questions or comments?

1-888-333-6792

*This product is not manufactured or distributed by McNEIL Consumer Healthcare, owner of the registered trademark Zyrtec®.


Drug Facts (continued)
■ 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 caplet once daily; do not take more than one 10 mg caplet 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)
Inactive Ingredients
colloidal silicon dioxide*, croscarmellose sodium*, hypromellose, lactose, magnesium stearate, maize starch*, microcrystalline cellulose*, polyethylene glycol, povidone*, titanium dioxide
*contains one or more of these ingredients
Questions or comments? 1-888-333-9792
LOT:
EXP:
ValuMeds
Compare to the active ingredient in Zyrtec®†
NDC 68210-0119-3
ALLERGY RELIEF
Cetirizine hydrochloride caplets, 10 mg / antihistamine
indoor & outdoor allergies
24 hour relief of:
• runny nose
• sneezing
• itchy, watery eyes
• itching of the nose or throat
30 caplets
Actual Size
†This product is not manufactured or distributed by McNeil Consumer Healthcare Division of McNEIL-PPC, Inc., owner of the registered trademark Zyrtec®.
Distributed By:
Spirit Pharmaceuticals, LLC
Ronkonkoma, NY 11779
ORIG 09/18
Made in India
ValuMeds
Compare to the active ingredient in Zyrtec®†
NDC 68210-0119-3
ALLERGY RELIEF
Cetirizine hydrochloride caplets, 10 mg / antihistamine
indoor & outdoor allergies
24 hour relief of:
• runny nose
• sneezing
• itchy, watery eyes
• itching of the nose or throat
30 caplets
Actual Size
KEEP CARTON FOR COMPLETE PRODUCT INFORMATION
Drug Facts
Active Ingredient (in each caplet) Purpose
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.
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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

CABINET ALLERGY RELIEF

cetirizine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68210-0119
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: 3WQ0SDW1A)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POVIDONE (UNII: FZ989GH94E)	

STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics			
Color	white (white to off-white)	**Score**	2 pieces
Shape	RECTANGLE (rounded-off rectangular biconvex tablet)	**Size**	10mm
Flavor		**Imprint Code**	G4
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210-0119-0	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/25/2019	
2	NDC:68210-0119-3	1 in 1 CARTON	03/10/2020	
2		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:68210-0119-1	300 in 1 BOTTLE; Type 0: Not a Combination Product	03/10/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA209274	10/25/2019	

Labeler - Spirit Pharmaceutical LLC (179621011)

Revised: 12/2024 Spirit Pharmaceutical LLC