ALL DAY ALLERGY- cetirizine hydrochloride capsule, liquid filled Rite Aid Corporation

Rite Aid Corporation All Day Allergy Drug Facts

Active ingredient (in each capsule)

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

and over	one 10 mg capsule once daily; do not take more than one 10 mg capsule 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	ask a doctor	
kidney disease		

Other information

- avoid high humidity and excessive heat above 40°C (104°F)
- protect from light

Inactive ingredients

butylated hydroxytoluene, gelatin, glycerin, mannitol, may contain pharmaceutical ink, polyethylene glycol 400, purified water, sodium hydroxide, sorbitan, sorbitol

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

Compare to the active ingredient of Zyrtec® Liquid Gels

ORIGINAL PRECRIPTION STRENGTH

24 HOUR

ALL DAY ALLERGY

CETIRIZINE HYDROCHLORIDE CAPSULES, 10 mg

ANTIHISTAMINE

INDOOR & OUTDOOR ALLERGIES

24-HOUR RELIEF OF

Sneezing

Runny nose

Itchy, watery eyes

Itchy throat or nose

ACTUAL SIZE

25 LIQUID GELS**

**LIQUID FILLED CAPSULES

NDC 11822-0980-1 Compare to the active ingredient of Zyrtec® Liquid Gels* **ORIGINAL** PRESCRIPTION STRENGTH **24 HOUR** L DAY **ALLERGY** YDROCHLORIDE CAPSULES, 10 mg ANTIHISTAMINE **INDOOR & OUTDOOR ALLERGIES** 24-HOUR RELIEF OF Sneezing Runny nose Itchy, watery eyes Itchy throat or nose **ACTUAL SIZE** .IQUID GELS**

**LIQUID FILLED CAPSULES

READ AND KEEP OUTER PACKAGING FOR COMPLETE WARNINGS AND INFORMATION

Store between 20-25°C (68-77°F).

DO NOT USE IF PRINTED FOIL UNDER CAP IS BROKEN OR MISSING

*This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., distributor of Zyrtec® Liquid Gels.

GLUTEN FREE

MADE IN CANADA

DISTRIBUTED BY:

RITE AID 30 HUNTER LANE CAMP HILL PA 17011 www.riteaid.com

SATISFACTION GUARANTEE

If you're not satisfied, we'll happily refund your money.



98N63 83 S1

Drug Facts

Active ingredient (in each capsule) Cetirizine HCl 10 mg......Antihistamine

Purpose

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 capsule once daily; do not take more than one 10 mg capsule 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 avoid high humidity and excessive heat above 40°C (104°F) ■ protect from light

Inactive ingredients butylated hydroxytoluene, gelatin, glycerin, mannitol, may contain pharmaceutical ink, polyethylene glycol 400, purified water, sodium hydroxide, sorbitan, sorbitol

Questions or comments?1-800-719-9260

ALL DAY ALLERGY

cetirizine hydrochloride capsule, liquid filled

Product Information

HUMAN OTC DRUG NDC:11822-0980 **Product Type** Item Code (Source)

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		
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)			
GELATIN, UNSPECIFIED (UNII: 2G86QN327L)			
GLYCERIN (UNII: PDC6A3C0OX)			
MANNITOL (UNII: 3OWL53L36A)			
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)			
WATER (UNII: 059QF0KO0R)			
SODIUM HYDROXIDE (UNII: 55X04QC32I)			
SORBITAN (UNII: 6O92ICV9RU)			
SORBITOL (UNII: 506T60A25R)			

Product Characteristics				
Color	YELLOW	Score	no score	
Shape	OVAL	Size	13mm	
Flavor		Imprint Code	C10	
Contains				

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822- 0980-1	1 in 1 PACKAGE	06/10/2022	
1		25 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11822- 0980-0	1 in 1 PACKAGE	06/10/2022	
2		40 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA213105	06/10/2022	

Labeler - Rite Aid Corporation (014578892)

Revised: 6/2022 Rite Aid Corporation