

ALLERGY RELIEF- cetirizine hydrochloride capsule, liquid filled
Rite Aid Corporation

Rite Aid Corporation Allergy Relief 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

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, 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 PRESCRIPTION STRENGTH

ALLERGY RELIEF

CETIRIZINE HYDROCHLORIDE CAPSULES, 10 mg

ANTIHISTAMINE

INDOOR & OUTDOOR ALLERGIES

24 HOUR RELIEF OF

Sneezing – Runny nose

Itchy, watery eyes

Itchy throat or nose

24 Hour

ACTUAL SIZE

40 LIQUID GELS**

**LIQUID FILLED CAPSULES

ORIGINAL PRESCRIPTION STRENGTH ALLERGY RELIEF

**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



40 LIQUID 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

Drug Facts

Active ingredient (in each capsule)	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.

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

SATISFACTION
GUARANTEE

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



011822999298

0Y349 83 S1

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, pharmaceutical ink, polyethylene glycol 400,
purified water, sodium hydroxide, sorbitan, sorbitol

Questions or comments? 1-800-719-9260

ALLERGY RELIEF

cetirizine hydrochloride capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-0457
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
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	10

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-0457-1	1 in 1 CARTON	11/30/2020	
1		40 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11822-0457-0	1 in 1 CARTON	12/03/2020	
2		25 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	ANDA207235	11/30/2020	

Labeler - Rite Aid Corporation (014578892)

Revised: 12/2020

Rite Aid Corporation