

**REXALL ALL DAY ALLERGY RELIEF- cetirizine hydrochloride tablet, film coated
Dolgenercorp, LLC**

Dolgenercorp, LLC All Day 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

- do not use if blister unit is broken or torn
- store between 20 to 25°C (68 to 77°F)

Inactive ingredients

corn starch, FD&C blue no. 1 aluminum lake, hypromellose, lactose monohydrate, magnesium stearate, polydextrose, polyethylene glycol, povidone, titanium dioxide, triacetin

Principal Display Panel

ORIGINAL PRESCRIPTION STRENGTH

All Day Allergy Relief

24 HOUR SYMPTOM RELIEF

CETIRIZINE HYDROCHLORIDE TABLETS 10 mg ANTIHISTAMINE

24 Hour relief of

- Sneezing
- Runny nose
- Itchy, watery eyes

- Itchy throat or nose

actual size

INDOOR & OUTDOOR ALLERGIES

14 Tablets

DISTRIBUTED BY OLD EAST MAIN CO.
100 MISSION RIDGE
GOODLETTSVILLE, TN 37072

Visit us at: Rexall.com
or call 1-866-4-REXALL



A0498



Since 1903

All Day Allergy Relief

CETIRIZINE HYDROCHLORIDE
TABLETS 10mg
ANTIHISTAMINE

ORIGINAL PRESCRIPTION STRENGTH



14 Tablets

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Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away. ▶	

TEAR ALONG PERFORATION, PEEL OFF PAPER AND PUSH PRODUCT THROUGH FOIL. IF DIFFICULT TO OPEN USE SCISSORS.

24 HOUR SYMPTOM RELIEF

4H266 CF C3



Since 1903

All Day Allergy Relief

CETIRIZINE HYDROCHLORIDE
TABLETS 10mg
ANTIHISTAMINE



14 Tablets



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All Day Allergy Relief

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24 Hour relief of

- Sneezing
- Itchy, watery eyes

actual size



• Runny nose • Itchy throat or nose

INDOOR & OUTDOOR ALLERGIES

14
Tablets

REXALL ALL DAY ALLERGY RELIEF

cetirizine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-699
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
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	WHITE	Score	no score
Shape	OVAL	Size	10mm
Flavor		Imprint Code	4H2
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-	1 in 1 CARTON	09/06/2012	

1	699-66	1 in 1 CARTON	09/06/2012	
1		14 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	ANDA078336	09/06/2012	

Labeler - Dolgencorp, LLC (068331990)

Revised: 6/2021

Dolgencorp, LLC