CHILDRENS ALL DAY ALLERGY- cetirizine hydrochloride solution Kroger Company

Children's All Day Allergy

Drug Facts

Active ingredient (in each 5 mL teaspoonful)

Cetirizine HCl 5 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- itchy, watery eyes
- sneezing
- 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

• use only with enclosed dosing cup

adults and children 6 years and over	take more than 2 teaspoonfuls (10 mL) once take more than 2 teaspoonfuls (10 mL) in 24 hours.		
adults 65 years and	1 teaspoonful (5 mL) once daily; do not take more than		
older	1 teaspoonful (5 mL) in 24 hours.		
children 2 to under 6 years of age	½ teaspoonful (2.5 mL) once daily. If needed, dose can be increased to a maximum of 1 teaspoonful (5 mL) once daily or ½ teaspoonful (2.5 mL) every 12 hours. Do not give more than 1 teaspoonful (5 mL) in 24 hours.		
children under 2 years of age	ask a doctor		
consumers with liver or kidney disease	ask a doctor		

Other information

- do not use if carton is opened, or if imprinted safety seal is broken or missing.
- see bottom panel for expiration date
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

bubble gum artificial flavor, glacial acetic acid, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium acetate anhydrous, sucralose

Questions?

1-800-632-6900

DISTRIBUTED BY THE KROGER CO. CINCINNATI, OHIO 45202

PRINCIPAL DISPLAY PANEL - 120 mL Bottle Carton

COMPARE TO the active ingredient in $CHILDREN'S\ ZYRTEC^{\ensuremath{\mathbb{R}}}$ *See side panel

NDC 30142-106-08

Kroger_®

FROM OUR FAMILY TO YOURS

2 YEARS & OLDER Children's All Day Allergy

Cetirizine Hydrochloride Oral Solution 1 mg/mL

Antihis tamine

SUGAR FREE

24 HOUR

24 Hour Relief of:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes Itchy Throat or Nose

• Our Pharmacists • Recommend

Indoor & Outdoor Allergies

Dosing Cup Included

Dye Free

Bubble Gum Flavor

4 FL OZ (120 mL)



Children's All Day Allergy

Cetirizine Hydrochloride Oral Solution 1 mg/mL **Antihistamine**

COMPARE TO the active ingredient in

□ COMPARE TO the active ingredient in □





2 YEARS & OLDER Children's All Day Allergy

Cetirizine Hydrochloride Oral Solution 1 mg/mL **Antihistamine**

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24 Hour Relief of:

- Sneezing
- Runny Nose
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24 Hour Relief of:

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- Runny Nose
- Itchy, Watery Eyes



NDC 30142-106-08







203 20581-0114-0

2 YEARS & OLDER Children's

Drug Facts (continued)

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Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.

Cetirizine Hydrochloride Oral Solution 1 mg/mL Antihistamine



DISTRIBUTED BY THE KROGER CO. CINCINNATI, OHIO 45202

QUALITY GUARANTEE www.kroger.com

MADE IN ISRAEL

*Children's Zyrtec is a registered trademark of UCB Pharma, S.A. CORPORATION BELGIUM.
UCB Pharma is not affiliated with The Kroger
Co. or this product.

Dosing cup should be washed and left to air dry after each use

SEALED WITH PRINTED NECKBAND

Drug Facts

Active ingredient (in each 5 mL teaspoonful) **Purpose**

Cetirizine HCl 5 mg.....Antihistamine

Uses temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

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- · itching of the nose or throat

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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 • use only with enclosed dosing cup		
adults and children 6 years and over	1 teaspoonful (5 mL) or 2 teaspoonfuls (10 mL) once daily depending upon severity of symptoms; do not take more than 2 teaspoonfuls (10 mL) in 24 hours.	
adults 65 years and older	1 teaspoonful (5 mL) once daily; do not take more than 1 teaspoonful (5 mL) in 24 hours.	
children 2 to under 6 years of age	½ teaspoonful (2.5 mL) once daily. If needed, dose can be increased to a maximum of 1 teaspoonful (5 mL) once daily or ½ teaspoonful (2.5 mL) every 12 hours. Do not give more than 1 teaspoonful (5 mL) in 24 hours.	
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Questions? 1-800-632-6900

CHILDRENS ALL DAY ALLERGY

cetirizine hydrochloride solution

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:30142-106

Route of Administration ORAL

Active Ingredient/Active Moiety

retive ingredient/retive wintery			
Ingredient Name	Basis of Strength	Strength	
Cetirizine Hydrochloride (UNII: 64O047KTOA) (Cetirizine - UNII:YO7261ME24)	Cetirizine Hydrochloride	5 mg in 5 mL	

Inactive Ingredients	
Ingredient Name	Strength
acetic acid (UNII: Q40Q9N063P)	
glycerin (UNII: PDC6A3C0OX)	
methylparaben (UNII: A2I8C7HI9T)	
propylene glycol (UNII: 6DC9Q167V3)	
propylparaben (UNII: Z8IX2SC1OH)	
water (UNII: 059QF0KO0R)	
sodium acetate anhydrous (UNII: NVG71ZZ7P0)	
sucralose (UNII: 96K6UQ3ZD4)	

Product Characteristics			
Color	YELLOW (colorless to slightly yellow)	Score	
Shape		Size	
Flavor	BUBBLE GUM (Sugar Free)	Imprint Code	
Contains			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:30142-106-08	1 in 1 CARTON		
1	120 mL in 1 BOTTLE		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA201546	05/20/2011	

Registrant - Taro Pharmaceuticals U.S.A., Inc. (145186370)

Establishment			
Name	Address	ID/FEI	Business Operations
Taro Pharmaceutical Industries Ltd.		600072078	MANUFACTURE(30142-106)

Revised: 4/2014 Kroger Company