

UP AND UP CHILDRENS ALLERGY RELIEF- cetirizine hcl solution
Target Corporation

Target Corporation Children's Allergy Relief Drug Facts

Active ingredient (in each 5 mL)

Cetirizine HCl 5 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

- use only with enclosed dosing cup
- find right dose on chart below
- mL = milliliter

adults and children 6 years and over	5 mL or 10 mL once daily depending upon severity of symptoms; do not take more than 10 mL in 24 hours.
adults 65 years and over	5 mL once daily; do not take more than 5 mL in 24 hours.
children 2 to under 6 years of age	2.5 mL once daily. If needed, dose can be increased to a maximum of 5 mL once daily or 2.5 mL every 12 hours. Do not give more than 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

- store between 20° to 25°C (68° to 77°F)
- do not use if carton is opened, or if printed neckband is broken or missing
- see bottom panel for lot number and expiration date

Inactive ingredients

anhydrous citric acid, artificial grape flavor, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose

Questions?

Call 1-888-547-7400

Package/Label Principal Display Panel

Compare to active ingredient in Children's Zyrtec®

children's

allergy relief

cetirizine hydrochloride oral solution 1 mg/mL

antihistamine

indoor and outdoor allergies

24 hour relief of:

sneezing

runny nose

itchy, watery eyes

itchy throat or nose

DYE AND SUGAR FREE

dosing cup included

24 HOUR

GRAPE FLAVOR

AGES 2+ YEARS

4 FL OZ (118 mL)



UP AND UP CHILDRENS ALLERGY RELIEF

cetirizine hcl solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-475
Route of Administration	ORAL		

Drug Facts (continued)

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GLUTEN FREE

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

C-000703-01-049
094 04 0295 R00 ID215427
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Minneapolis, MN 55403
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*This product is not manufactured or distributed by McNeil Consumer Healthcare, division of McNeil-PPC, Inc., distributor of Children's Zyrtec®.

children's allergy relief
cetirizine hydrochloride oral solution 1 mg/mL antihistamine

100% satisfaction guaranteed or your money back.

Each year we give 5% of our profit to communities, which adds up to more than \$4 Million a week. See all the good we do together at Target.com/Community.

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3 70030 62996 1

LOT NO.

EXP.

: 47526 UV C1

Active Ingredient/Active Moiety

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
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	GRAPE	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-475-34	1 in 1 CARTON	10/15/2015	
1		236 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:11673-475-26	1 in 1 CARTON	01/19/2018	
2		118 mL 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	ANDA204226	10/15/2015	

Labeler - Target Corporation (006961700)

Revised: 1/2018

Target Corporation