CHILDRENS CETIRIZINE HYDROCHLORIDE- cetirizine hcl solution ATLANTIC BIOLOGICALS CORP.

Major Pharmaceuticals Children's Cetirizine Hydrochloride Oral Solution 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 or comments?

1-800-616-2471

DISTRIBUTED BY:

ATLANTIC BIOLOGICALS CORP.

MIAMI, FL 33179

Package/Label Principal Display Panel

Children's Cetirizine Hydrochloride Oral Solution 1 mg/mL

Antihistamine

ALLERGY

INDOOR & OUTDOOR ALLERGIES

24 HOUR RELIEF OF:

Sneezing / Runny Nose

Itchy, Watery Eyes / Itchy Throat or Nose

2yrs & older

Dosing Cup Included

Dye-Free

Sugar-Free

Grape Flavored Syrup

CHILDREN'S
CETIRIZINE HCL
ORAL SOLUTION
DELIVERS 5mg/5mL
GRAPE FLAVOR
DYE/GLUTEN/SUGAR FREE

EXP: XX/XX/XX LOT: XXXXXX

DIST. BY:

ATLANTIC BIOLOGICALS CORP.

17856676501 REV.1

17856-6765-01 CHILDREN'S CETIRIZINE HCL ORAL SOLUTION 1mg PER 1mL DELIVERS 5mg/5mL



See package insert for indications and dosage schedule

Dye free, Gluten Free, Sugar Free Grape Flavored Antihistamine for 24 hr relief of allergies. Store between 20° to 25°C (68° to 77°F).

Keep this and all medications out of the reach of children



17856-6765-01 Dosage 5mg/5mL

CHILDREN'S CETIRIZINE HCL ORAL SOLUTION

Qty: 72 CUPS

GTIN: 00317856676519

Exp: 12/12/24

Lot: XXXXXXXXXX

OTC

Packaged by:

Distributed by: Atlantic Biologicals Corp. Miami, FL 33179

Rev.08/21

Call to Reorder:

CHILDRENS CETIRIZINE HYDROCHLORIDE

cetirizine hcl solution

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:17856-6765(NDC:0904-6765)
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZ INE 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	
NDC:17856- 6765-1	72 in 1 BOX, UNIT-DOSE	12/12/2024		
1 NDC:17856- 6765-2	5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204226	08/22/2018	

Labeler - ATLANTIC BIOLOGICALS CORP. (047437707)

Establishment				
Name	Address	ID/FEI	Business Operations	
UNIT DOSE SOLUTION		360804194	repack(17856-6765)	

Revised: 12/2024 ATLANTIC BIOLOGICALS CORP.