

CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride solution
STRATEGIC SOURCING SERVICES LLC

Cetirizine Hydrochloride

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

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

Inactive ingredients

artificial grape flavor, glacial acetic acid, glycerin, methylparaben, natural and artificial banana flavor, propylene glycol, propylparaben, purified water, sodium acetate (anhydrous), sucralose

Questions?

call **833-358-6431** Monday to Friday 9:00am to 7:00pm EST

Distributed by: McKesson Corp.,
via Strategic Sourcing Services LLC.
Memphis, TN 38141

PRINCIPAL DISPLAY PANEL - 120 mL Bottle Carton

NDC 70677-1236-1

COMPARE TO THE
ACTIVE INGREDIENT
IN CHILDREN'S ZYRTEC®*

Foster & Thrive™

Children's
All Day
Allergy

Cetirizine Hydrochloride
Oral Solution
1 mg/1 mL
ANTIHISTAMINE Allergy

Indoor & Outdoor Allergies
24 hour relief of:

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

Sugar Free

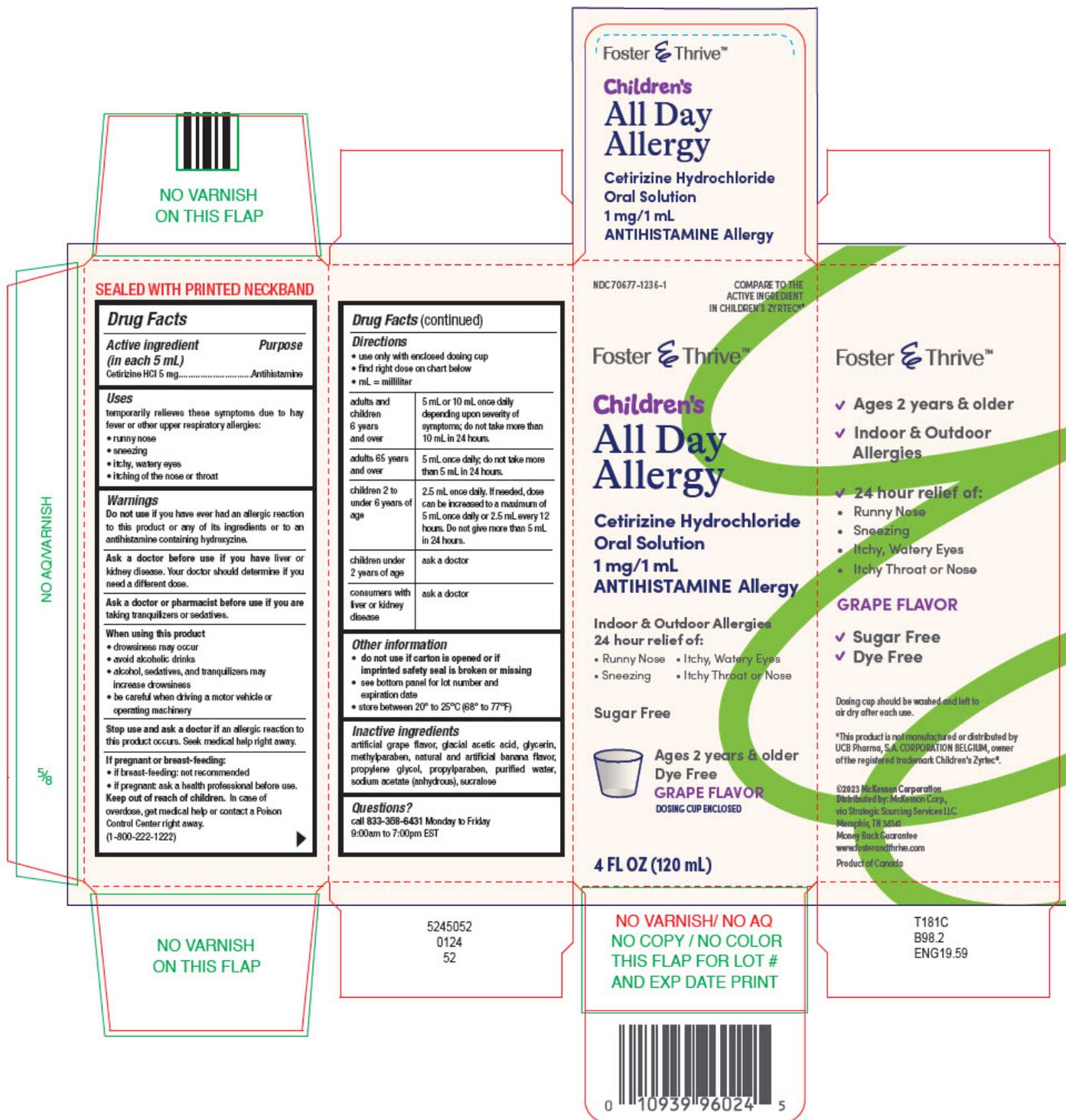
Ages 2 years & older

Dye Free

GRAPE FLAVOR

DOSING CUP ENCLOSED

4 FL OZ (120 mL)



CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-1236
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name			Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)			CETIRIZINE HYDROCHLORIDE	1 mg in 1 mL
Inactive Ingredients				
Ingredient Name			Strength	
ACETIC ACID (UNII: Q40Q9N063P)				
GLYCERIN (UNII: PDC6A3C00X)				
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	GRAPE		Imprint Code	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70677-1236-1	1 in 1 CARTON	02/02/2024	
1		120 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	ANDA090182	02/02/2024		

Labeler - STRATEGIC SOURCING SERVICES LLC (116956644)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(70677-1236)