

TOPCARE CHILDRENS ALL DAY ALLERGY- cetirizine hcl solution
Topco Associates LLC

Topco Associates LLC. Children's All-Day Allergy 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 bubble gum flavor, propylene glycol, purified water, sodium benzoate, sorbitol solution, sucralose

Questions or comments?

1-888-423-0139

Package/Label Principal Display Panel

COMPARE TO CHILDREN'S ZYRTEC® ACTIVE INGREDIENT

children's

All-Day Allergy

CETIRIZINE HYDROCHLORIDE ORAL SOLUTION 1 mg / mL

ANTIHISTAMINE

OUR PHARMACISTS RECOMMEND

INDOOR & OUTDOOR ALLERGIES

24 HOUR

RELIEF OF:

Sneezing

Runny Nose

Itchy, Watery Eyes

Itchy Throat or Nose

Dye Free & Sugar Free

2 Years & older

BUBBLE GUM FLAVOR

4 FL OZ (118 mL)

Dosing cup included



TOPCARE CHILDRENS ALL DAY ALLERGY

cetirizine hcl solution

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:36800-189 |
| Route of Administration | ORAL | | |

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 | YELLOW (Clear to light yellow) | Score | |
| Shape | | Size | |
| Flavor | BUBBLE GUM | Imprint Code | |
| Contains | | | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:36800-189-26 | 1 in 1 CARTON | 12/23/2013 | |
| 1 | | 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 | 12/23/2013 | |

Labeler - Topco Associates LLC (006935977)

Revised: 11/2019

Topco Associates LLC