# OUR FAMILY CHILDRENS CETIRIZINE- cetirizine hydrochloride solution Nash Finch Company

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Our Family<sup>®</sup> Childrens Cetirizine

**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
- 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

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 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.
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)

## **Inactive ingredients**

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

## Questions?

Call **1-866-923-4914** 

DISTRIBUTED BY

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MINNEAPOLIS, MN 55435

## PRINCIPAL DISPLAY PANEL - 120 mL Bottle Carton

**Oral Solution** 

Compare to the active ingredient in Children's Zyrtec®\*

NDC 70253-106-08

Our Family®

Quality Care Since 1904

Children's
CETIRIZINE
HYDROCHLORIDE
ORAL SOLUTION
1 mg/mL Antihis tamine

**SUGAR FREE & DYE FREE** 

**INDOOR & OUTDOOR ALLERGIES** 

24 hour relief of:

Runny noseSneezing

Itchy, watery eyes Itchy throat or nose

2 Yrs. & older

Dosing Cup Included

**Bubble Gum** Flavor

4 FL OZ (120 mL)

# **Oral Solution**

# Children's YDROCHLORIDE **ORAL SOLUTION** 1 mg/mL Antihistamine SUGAR FREE & DYE FREE

NDC 70253-106-08

**Oral Solution** 

**Oral Solution** 

NDC 70253-106-08



# Children's HYDROCHLORIDE ORAL SOLUTION

1 mg/mL Antihistamine

SUGAR FREE & DYE FREE

INDOOR & OUTDOOR ALLERGIES

## 24 hour relief of:

- Runny nose
   Itchy, watery eyes
- Sneezing
- Itchy throat or nose



# Children's CETIRIZINE **HYDROCHLORIDE ORAL SOLUTION**

1 mg/mL Antihistamine

SUGAR FREE & DYE FREE

INDOOR & OUTDOOR ALLERGIES

# 24 hour relief of:

- Runny nose
- Itchy, watery eyes
- Sneezing
- · Itchy throat or nose

2 Yrs. & older



Dosing Cup Induded

FL OZ

2 Yrs. & older



Dosing Cup Induded





0-2150-62202

# Children's CETIRIZINE HYDROCHLORIDE ORAL SOLUTION 1 mg/mL Antihistamine

SUGAR FREE & DYE FREE

# 24 hour relief of:

Runny nose Sneezing

- Itchy, watery eyes
- · Itchy throat or nose

Dosing cup should be washed and left to air dry after each use. Do not use if carton is opened, or if imprinted safety seal is broken or missing. See bottom panel for expiration date.



Dosing Cup Included

## 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.

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			
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		

adults 65 years and older

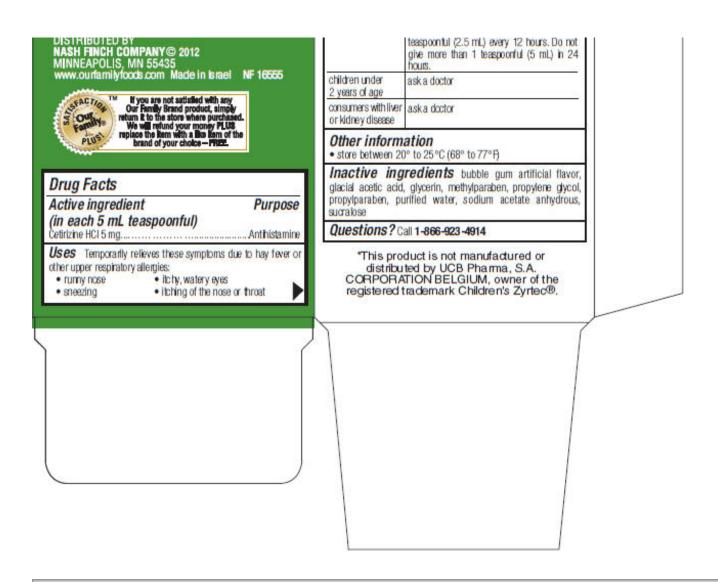
1 teaspoonful (5 mL) once daily; do not take more than 1 teaspoonful (5 mL) in 24 hours.

teaspoonfuls (10 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

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## **OUR FAMILY CHILDRENS CETIRIZINE**

cetirizine hydrochloride solution

<b>Product Information</b>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70253-106
Route of Administration	ORAL		

Active Ingredient/Active Moiety				
Ingredient Name Basis of Strength Strength				
Cetirizine Hydrochloride (UNII: 640047KTOA) (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			

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70253-106-08	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	

## **Labeler** - Nash Finch Company (006962294)

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

Establishment				
Name	Address	ID/FEI	Business Operations	
Taro Pharmaceutical Industries, Ltd.		600072078	MANUFACTURE(70253-106)	

Revised: 12/2012 Nash Finch Company