

CHILDRENS LORATADINE SUGAR FREE- loratadine solution
Sun Pharmaceutical Industries, Inc.

Childrens Loratadine Oral Solution
Sugar Free

Drug Facts

Active ingredient (in each 5 mL teaspoonful)

Loratadine 5 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- itchy, watery eyes
- sneezing
- 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.

Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding, 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 | 2 teaspoonfuls (tsp) daily; do not take

adults and children 10 years and over	more than 2 teaspoonfuls (tsp) in 24 hours
children 2 to under 6 years of age	1 teaspoonful (tsp) daily; do not take more than 1 teaspoonful (tsp) 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 bottle wrap imprinted with "SEALED FOR SAFETY" is broken or missing.**
- see bottom panel for lot number and expiration date
- store between 20° and 25°C (68° and 77°F)

Inactive ingredients

butylated hydroxyanisole, glycerin, grape flavor, maltitol solution, masking agent, phosphoric acid, polyethylene glycol, propylene glycol, purified water, sodium benzoate, sodium phosphate monobasic dihydrate, sorbitol solution, sucralose powder.

Questions?

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

Distributed by:

Taro Pharmaceuticals U.S.A., Inc.

Hawthorne, NY 10532

PRINCIPAL DISPLAY PANEL - 120 mL Bottle Carton

NDC 51672-2131-8

Compare to the active ingredient in Children's Claritin[®]*

Original Prescription Strength

Children's

Loratadine

Oral Solution

USP, 5 mg/5 mL

(Antihistamine) **ALLERGY**

Non-Drowsy[†]

Indoor & Outdoor Allergies

SUGAR FREE

24

Hour

Relief of:

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

Ages 2 years and older

Dosing Cup

Enclosed

**Grape
Flavor**

†When taken as directed.
See Drug Facts Panel.

4 FL OZ
(120 mL)

NDC 51672-2131-8

Compare to the active ingredient in Children's Claritin®*

Children's Loratadine Oral Solution USP, 5 mg/5 mL

(Antihistamine) **ALLERGY**

Non-Drowsy†

Indoor & Outdoor Allergies

NDC 51672-2131-8

Compare to the active ingredient in Children's Claritin®*

Original Prescription Strength

Children's Loratadine Oral Solution USP, 5 mg/5 mL

(Antihistamine) **ALLERGY**

Non-Drowsy†

Indoor & Outdoor Allergies

SUGAR FREE

24 Hour

Relief of:

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

Ages 2 years and older

Dosing Cup Enclosed



Grape Flavor

4 FL OZ

†When taken as directed.

Original Prescription Strength

Children's Loratadine Oral Solution USP, 5 mg/5 mL

(Antihistamine) **ALLERGY**

Non-Drowsy†

Indoor & Outdoor Allergies

- Alcohol Free
- Sugar Free
- 24 Hour Relief
- Dye Free

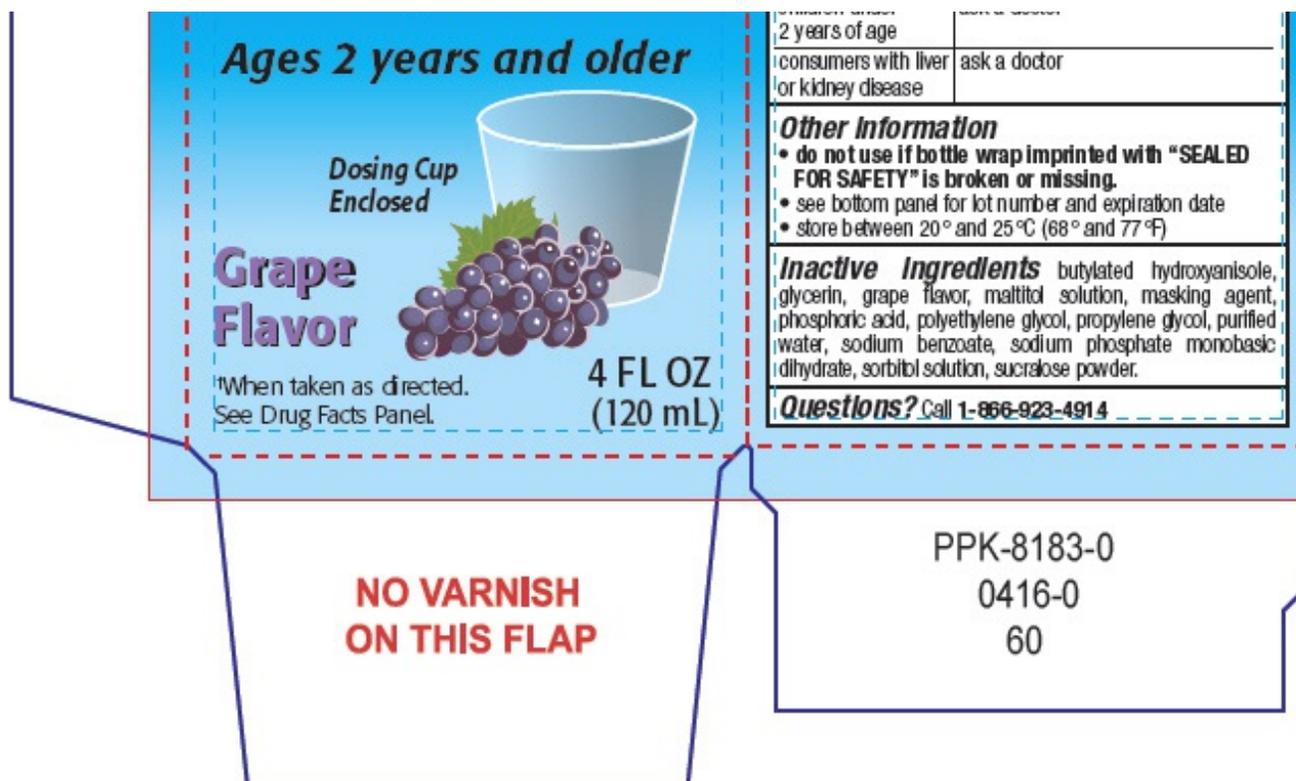
Dosing Cup Enclosed



Grape Flavor

*All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Children's Claritin®.

Distributed by:
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Hawthorne, NY 10532
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CHILDRENS LORATADINE SUGAR FREE

loratadine solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51672-2131
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII: 7AJ03BO7QN)	LORATADINE	5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
BUTYLATED HYDROXYANISOLE (UNII: REK4960K2U)	
GLYCERIN (UNII: PDC6A3C0OX)	
SORBITOL (UNII: 506T60A25R)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM PHOSPHATE, MONOBASIC, DIHYDRATE (UNII: 5QWK665956)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
MALTITOL (UNII: D65DG142WK)	

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:51672-2131-4	1 in 1 CARTON	03/17/2017	
1		60 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:51672-2131-8	1 in 1 CARTON	03/17/2017	
2		120 mL in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:51672-2131-1	1 in 1 CARTON	03/17/2017	
3		240 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	ANDA076805	03/17/2017	

Labeler - Sun Pharmaceutical Industries, Inc. (146974886)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(51672-2131)

Revised: 7/2025

Sun Pharmaceutical Industries, Inc.