

LORATADINE- loratadine solution
Cardinal Health

Loratadine Oral Solution, 5 mg/5 mL

Drug Facts

Active ingredient (in each 5 mL)

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.

Directions

adults and children 6 years and over	2 teaspoonfuls daily; do not take more than 2 teaspoonfuls in 24 hours
children 2 to under 6 years of age	1 teaspoonful daily; do not take more than 1 teaspoonful in 24 hours
consumers with liver or kidney disease	ask a doctor

Other information

- **safety sealed: do not use if imprinted safety seal is torn or missing**

- store between 2° and 25°C (36° and 77°F)

Inactive ingredients

artificial peach flavor, citric acid monohydrate, glycerin, propylene glycol, purified water, sodium benzoate, sodium metabisulfite, sucrose

Questions?

Call 1-800-200-6313

DISTRIBUTED BY
CARDINAL HEALTH
DUBLIN, OHIO 43017

PRINCIPAL DISPLAY PANEL - 120 mL Bottle Carton

LEADER®

NDC 37205-378-26

*Compare to
Children's Claritin®
active ingredient†*

*Original Prescription
Strength*

NON-DROWSY*

**Allergy
Relief**

*Loratadine Syrup
(Loratadine Oral Solution) 5 mg/5 mL*

Antihistamine

24 Hour

*Relief of: Sneezing, Runny Nose,
Itchy, Watery Eyes • Itchy Throat or Nose*

Fruit Flavor

**When taken as directed.
See Drug Facts Panel.*

**Ages two years
and older**

**SATISFACTION
GUARANTEED**

4 FL OZ (120 mL)

478
70506-1109-2



NDC 37205-378-26

Original Prescription
Strength

Compare to
Children's Claritin®
active ingredient

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SEALED WITH PRINTED NECKBAND

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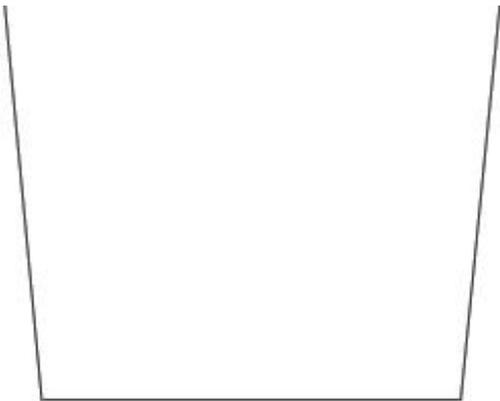
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Allergy Relief

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Antihistamine

24Hour



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NDC 37205-378-26

Original Prescription Strength

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

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LORATADINE

loratadine solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37205-378
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
citric acid monohydrate (UNII: 2968PHW8QP)	
glycerin (UNII: PDC6A3C0OX)	
propylene glycol (UNII: 6DC9Q167V3)	
water (UNII: 059QF0KO0R)	

sodium benzoate (UNII: OJ245FE5EU)	
sodium metabisulfite (UNII: 4VON5FNS3C)	
sucrose (UNII: C151H8M554)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	PEACH	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37205-378-26	1 in 1 CARTON	08/20/2004	
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	ANDA076805	08/20/2004	

Labeler - Cardinal Health (097537435)

Establishment

Name	Address	ID/FEI	Business Operations
Taro Pharmaceutical Industries Ltd.		600072078	MANUFACTURE(37205-378)

Revised: 12/2019

Cardinal Health