

SUNMARK CHILDRENS LORATADINE- loratadine solution
Strategic Sourcing Services LLC

Sunmark®
children's loratadine syrup

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	2 teaspoonfuls daily; do not take
children 6 years	

and over	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 1-866-923-4914

Distributed by McKesson
One Post Street, San Francisco, CA 94104

PRINCIPAL DISPLAY PANEL - 120 mL Bottle Carton

sunmark®

COMPARE TO
CHILDREN'S CLARITIN®
ACTIVE INGREDIENT*

NDC 49348-636-34

**24 HOUR
ALLERGY RELIEF**

**children's
loratadine
syrup**

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

Relief of
sneezing, runny nose
itchy, watery eyes
itchy throat or nose

Dye Free

Non-drowsy†
Ages two years & older

FRUIT FLAVOR

4 FL OZ (120 mL)

†When taken as directed. See Drug Facts Panel.



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Loratadine 5 mg	Antihistamine

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- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

Warnings
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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

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*This product is not manufactured or distributed by Schering-Plough HealthCare Products Inc., owner of the registered trademark Claritin®.

McKesson
Empowering healthcare

Another Quality Product
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One Post Street, San Francisco, CA 94104
Money Back Guarantee
Please visit us at www.sunmarkbrand.com
MADE IN ISRAEL.



SUNMARK CHILDRENS LORATADINE

loratadine solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49348-636
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	yellow (colorless to slightly yellow)	Score	
Shape		Size	
Flavor	FRUIT	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49348-636-34	1 in 1 CARTON	11/14/2012	
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 - Strategic Sourcing Services LLC (116956644)

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

Establishment

Name	Address	ID/FEI	Business Operations
Taro Pharmaceutical Industries Ltd.		600072078	manufacture(49348-636)

Revised: 12/2024

Strategic Sourcing Services LLC