

SUNMARK CHILDRENS LORATADINE- loratadine solution
McKesson

sunmark[®]
Children's Loratadine

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 years and over	2 teaspoonfuls (tsp) daily; do not take 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

- **safety sealed: do not use if imprinted safety seal is torn or missing**
- store between 20° and 25°C (68° and 77°F)

Inactive ingredients

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

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 49348-333-34

**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

SUGAR FREE
GRAPE FLAVOR

4 FL OZ (120 mL)

†When taken as directed. See Drug Facts Panel.



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- Alcohol Free
- Sugar Free
- 24 Hour Relief
- Dosage Cup Enclosed

* This product is not manufactured or distributed by MSD Consumer Care Inc., a subsidiary of Merck & Co., Inc.

MEKESON



GRAPE FLAVOR

4 FL OZ (120 mL)

† When taken as directed. See Drug Facts Panel.

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

**NO COPY ON THIS FLAP
FOR LOT # AND EXPIRY
DATE PRINT**

T181B



**NO VARNISH
ON THIS FLAP**



COMPARE TO
CHILDREN'S CLARITIN®
ACTIVE INGREDIENT*
NDC 49348-333-34

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children's
loratadine

SEALED WITH PRINTED NECKBAND

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**NO VARNISH
ON THIS FLAP**

PPK-7109-1
0913-1
M166

SUNMARK CHILDRENS LORATADINE

loratadine solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49348-333
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
glycerin (UNII: PDC6A3C0OX)	
sorbitol (UNII: 506T60A25R)	
phosphoric acid (UNII: E4GA8884NN)	
polyethylene glycols (UNII: 3WJQ0SDW1A)	
propylene glycol (UNII: 6DC9Q167V3)	
water (UNII: 059QF0KO0R)	
sodium benzoate (UNII: OJ245FE5EU)	
sodium metabisulfite (UNII: 4VON5FNS3C)	
sodium phosphate, monobasic, dihydrate (UNII: 5QWK665956)	

Product Characteristics

Color	YELLOW (colorless to slightly yellow)	Score	
Shape		Size	
Flavor	GRAPE (sugar free)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49348-333-34	1 in 1 CARTON		
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	02/27/2010	

Labeler - McKesson (177667227)

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