

ALL DAY ALLERGY RELIEF- loratadine tablet
Safeway, Inc.

Drug Facts

Active ingredient (in each tablet)

Loratadine, USP 10 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.

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 (1-800-222-1222) right away.

Directions

adults and children 6 years and over	1 tablet daily; not more than 1 tablet in 24 hours
children under 6 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- store at 20°-25°C (68°-77°F) (see USP Controlled Room Temperature)
- protect from light

Inactive ingredients

lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

COMPARE TO Claritin® 24 Hour active ingredient†

All Day Allergy Relief

LORATADINE TABLETS, 10mg

Antihistamine

Non-Drowsy*

Indoor & Outdoor Allergies

24 hour relief of:

- Sneezing
- Runny Nose
- Itchy, watery eyes
- Itchy Throat or Nose

TABLETS

*When taken as directed. See Drug Facts panel.

*This product is not manufactured or distributed by Bayer Healthcare LLC, distributor of Claritin® 24 Hour.

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

DISTRIBUTED BY:

BETTER LIVING BRANDS LLC

P.O. BOX 99, PLEASANTON, CA 94566-0009

Package Label

Drug Facts Active ingredient (in each tablet) Loratadine, USP 10 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. 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 (1-800-222-1222) right away.							
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NDC 21130-914-60

All Day Allergy Relief
 LORATADINE TABLETS, 10 mg
 Antihistamine

Non-drowsy*
Indoor & outdoor allergies

24-hour relief of

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

Actual Size

60 TABLETS

*When taken as directed. See Drug Facts panel.

NDC 21130-914-60

COMPARE TO
 Claritin® 24 Hour
 active ingredient!
 NDC 21130-914-60

All Day Allergy Relief
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PLD-A604A FC008469 Product of India

Lot No.:
Exp. Date:

Scan here for more information

S2801 RD 23047

 3 21130 78897 3
1 BOTTLE INSIDE

SIGNATURE CARE All Day Allergy Relief

ALL DAY ALLERGY RELIEF			
loratadine tablet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-914
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg	

Inactive Ingredients

Ingredient Name	Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	439
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-914-10	10 in 1 CARTON	03/31/2023	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:21130-914-60	1 in 1 BOX	03/31/2023	
2		60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:21130-914-12	1 in 1 BOX	03/31/2023	
3		120 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075209	03/31/2023	

Labeler - Safeway, Inc. (009137209)

Revised: 5/2023

Safeway, Inc.