

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

Gluten-free

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							
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Signature SELECT
ALL DAY ALLERGY RELIEF
LORATADINE 10 mg TABLETS
ANTIHISTAMINE

Non-drowsy*
Indoor & Outdoor Allergies

24-hour relief of

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

Gluten-free
*When taken as directed. See Drug Facts panel.

60 Tablets

COMPARE TO
Claritin® 24 Hour
Active Ingredient!
NDC 21130-914-60

NDC 21130-914-60

Signature SELECT
ALL DAY ALLERGY RELIEF
LORATADINE 10 mg TABLETS
ANTIHISTAMINE

Non-drowsy*
Indoor & Outdoor Allergies

24-hour relief of

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

Actual Size

Gluten-free
*When taken as directed. See Drug Facts panel.

60 Tablets

LOVE THE PLANET

DISTRIBUTED BY:
 BETTER LIVING BRANDS LLC
 P.O. BOX 99
 PLEASANTON, CA 94566-0009
 †1-888-723-3929

nonrecycle.info

PAPER BOX PLASTIC BOTTLE

Product of India
 PLD-B604A FCC09072

Lot No:
 Exp. Date:

Scan here for more information

RD24226 S2801
3 21130 478897 3

1 BOTTLE INSIDE

SIGNATURE SELECT 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	03/31/2027
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:21130-914-60	1 in 1 BOX	03/31/2023	03/31/2027
2		60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:21130-914-12	1 in 1 BOX	03/31/2023	03/31/2027
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	03/31/2027

Labeler - Safeway, Inc. (009137209)

Revised: 1/2026

Safeway, Inc.