

LORATADINE- loratadine tablet
Safecor Health, LLC

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
- 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	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

- **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.**
- Store at 20° to 25°C (68° to 77°F). [See USP Controlled Room Temperature.]
- protect from excessive moisture

Inactive ingredients

Corn starch, lactose monohydrate and magnesium stearate.

Questions or comments?

call **1-800-447-1006**

- Serious side effects associated with use of this product may be reported to this number.

Manufactured for:

Mylan Pharmaceuticals Inc.
Morgantown, WV 26505 U.S.A.

Made in India

Code No.: MH/DRUGS/25/NKD/89

Distributed by:

Safecor Health LLC
Rockford, IL 61103 U.S.A.

PRODUCT PACKAGING

NDC 48433-064-20

Loratadine
Tablets USP, 10 mg

Antihistamine

Original Prescription Strength Non-Drowsy*

Indoor and Outdoor Allergies

24 Hour Relief(See Uses section of Patient Information)

***When taken as directed. See Patient Information.**

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Loratadine **10 mg**
Tablets, USP

Antihistamine **Non-Drowsy***
Original Prescription Strength
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24 Hour Relief (See Uses section of Patient Information)
*When taken as directed. See Patient Information.
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NDC 48433-064-20

Loratadine **10 mg**
Tablets, USP

Antihistamine



100 Tablets (10 x 10)

Drug Facts

Active ingredient (in each tablet)	Purpose
Loratadine USP, 10 mg	Antihistamine

Uses Scan QR code for Patient Information


Warnings Scan QR code for Patient Information

Directions Scan QR code for Patient Information

Other information ■ **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.** ■ Store at 20° to 25°C (68° to 77°F). [See USP Controlled Room Temperature.] ■ protect from excessive moisture

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Manufactured for:
Mylan Pharmaceuticals Inc.
Morgantown, WV 26505 U.S.A.
Made in India

Scan for current Patient Information and Prescribing Information. 

Code No.: MWDRUS/25/NND/89 S-13206



- This unit dose package is not child resistant.
- For institutional use only.
- Keep this and all drugs out of the reach of children.
- This container provides light-resistance.
- See window for lot number and expiration date.

Distributed by:
Safecor Health LLC, Rockford, IL 61103 U.S.A.

1-800-447-1006 SafecorHealth.com

LORATADINE

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48433-064(NDC:0378-8880)
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
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	white (white to off-white)	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	G;L;10

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48433-064-20	100 in 1 CARTON	03/24/2026	
1	NDC:48433-064-01	1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076154	03/24/2026	

Labeler - Safecor Health, LLC (828269675)

Revised: 3/2026

Safecor Health, LLC