LORATADINE ALLERGY RELIEF- loratadine tablet
OHM LABORATORIES INC.

Loratadine Allergy Relief

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
- 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

adults and children 6 years and over	1 tablet daily; not more than 1 tablet in 24 hours
children under 6 years of	ask a doctor

age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other Information

- store between 20° to 25° C (68° to 77° F)
- protect from excessive moisture
- **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING**.

Inactive ingredients

corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

Questions?

call **1-800-406-7984**

Distributed by:
Ohm Laboratories Inc.
New Brunswick, NJ 08901

PRINCIPAL DISPLAY PANEL - 10 mg Tablet Blister Pack Carton

†Compare To
the active ingredient of
Claritin®

ohm®

NDC 51660-526-11

NON-DROWSY*
24 Hour
Allergy Relief

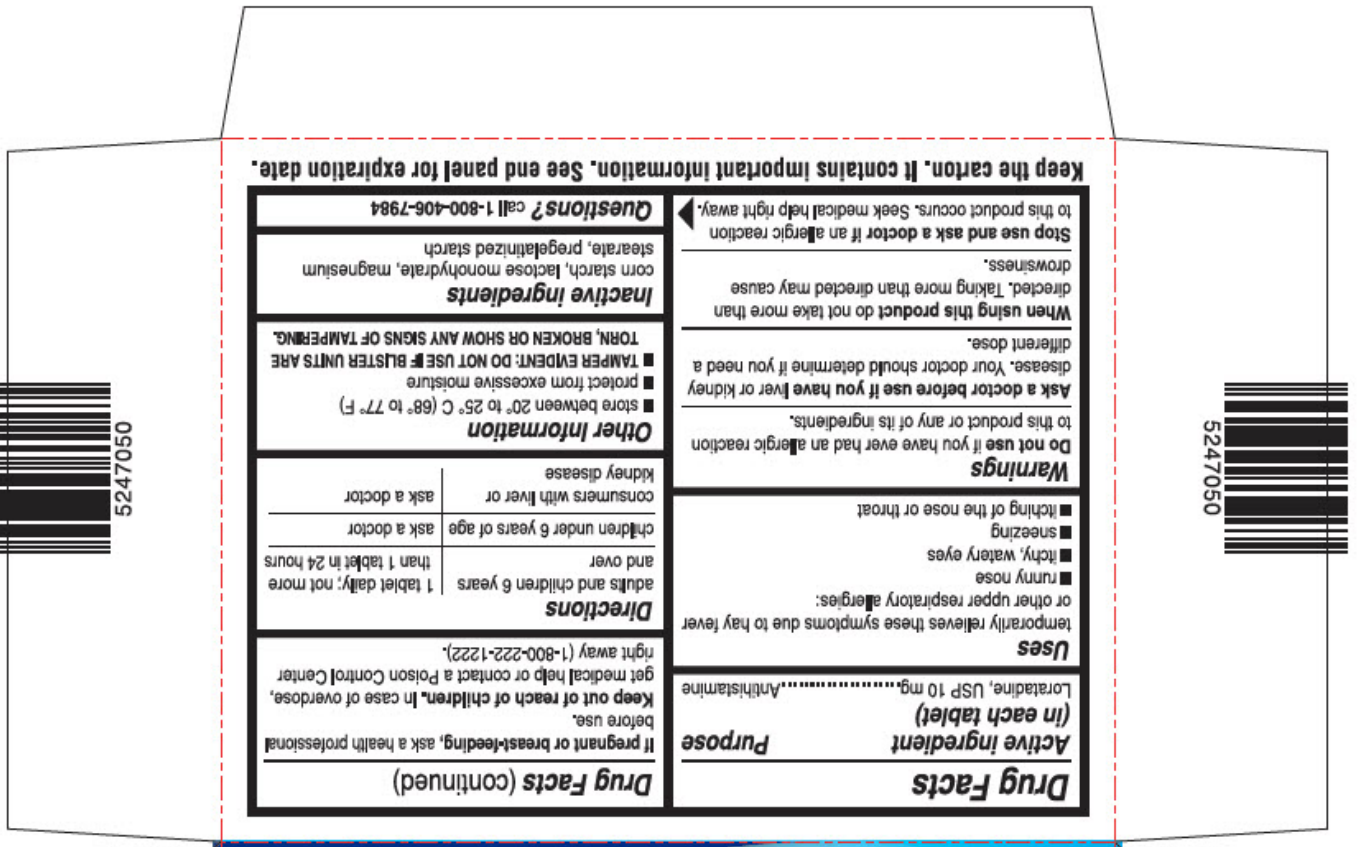
Original Prescription Strength
Allergy Relief
Loratadine Tablets, USP 10 mg
Antihistamine
Indoor & Outdoor Allergies

Relief of:

- ✓ Sneezing
- ✓ Runny Nose
- ✓ Itchy, Watery Eyes
- ✓ Itchy Throat or Nose

10 Tablets

* When taken as directed. See Drug Facts Panel.



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Drug Facts (continued)

Keep the carton. It contains important information. See end panel for expiration date.

5247050

5247050

ohm®

Allergy Relief

Loratadine Tablets, USP 10 mg

Antihistamine

Indoor & Outdoor Allergies

NDC 51660-526-11

†Compare To the active ingredient of Claritin®

ohm®

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24 Hour Allergy Relief

Original Prescription Strength

Allergy Relief

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Antihistamine

Indoor & Outdoor Allergies

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R1123

Distributed by:
Ohm Laboratories Inc.
New Brunswick, NJ 08901

†Ohm is a registered trademark of Sun Pharmaceutical Industries, Inc.
All other trademarks are property of their respective owners.

Batch No.

Expiration Date:

Non Varnish Area

ohm®

Allergy Relief

Loratadine Tablets, USP 10 mg

Antihistamine

LORATADINE ALLERGY RELIEF

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51660-526
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	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**	RX526
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51660-526-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2017	
2	NDC:51660-526-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2017	
3	NDC:51660-526-53	300 in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2017	
4	NDC:51660-526-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2019	
5	NDC:51660-526-31	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	11/01/2017	
6	NDC:51660-526-11	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	11/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076134	11/01/2017	

Labeler - OHM LABORATORIES INC. (184769029)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	MANUFACTURE(51660-526)

Revised: 12/2023 OHM LABORATORIES INC.