LORATADINE- loratadine tablet
Ohm Laboratories 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
- 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 age	ask a doctor
consumers with liver or kidney disease	ask a doctor

OTHER INFORMATION

- store between 20° and 25° C (68° and 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**

PRINCIPAL DISPLAY PANEL

SOUND BODY™

COMPARE TO THE ACTIVE INGREDIENT IN CLARITIN®†

Relief of: Sneezing, Itchy, Watery Eyes; Runny Nose; Itchy Throat or Nose

***When taken as directed. See Drug Facts Panel.**

Original Prescription Strength

NON-DROWSY*

Allergy Relief

Loratadine Tablets, USP 10 mg

Antihistamine

Indoor & Outdoor Allergies

10 Tablets

24 HOURS

ALLERGY RELIEF

Manufactured by: Ohm Laboratories Inc.

5104668/R0713

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

Drug Facts

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Loratadine, USP 10 mg	Antihistamine

Uses

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Drug Facts (continued)

If pregnant or breast-feeding, ask a health professional before use.

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Inactive ingredients

corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

Questions?

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

5104668

5104668

SOUND BODY™

Allergy Relief

Loratadine Tablets, USP 10 mg

Antihistamine
Indoor & Outdoor Allergies

SOUND BODY™

COMPARE TO THE ACTIVE INGREDIENT IN **CLARITIN®†**

Actual Size

Relief of: Sneezing; Itchy, Watery Eyes; Runny Nose; Itchy Throat or Nose

* When taken as directed. See Drug Facts Panel.

Original Prescription Strength
NON-DROWSY*

Allergy Relief

Loratadine Tablets, USP 10 mg

Antihistamine
Indoor & Outdoor Allergies

10 Tablets

24 HOURS
Allergy Relief

†All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Claritin®.

Batch No.

Expiration Date:

Non Varnish Area

R0713

Manufactured by:
Ohm Laboratories Inc.
1385 Livingston Avenue
North Brunswick, NJ 08902

V#1008633 ITEM#1090311

N 3 51660 52669 1

SOUND BODY™

Allergy Relief

Loratadine Tablets, USP 10 mg

Antihistamine
Indoor & Outdoor Allergies

LORATADINE

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51660-209
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**	RX;526
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:51660-209-69	1 in 1 CARTON		
1		10 in 1 BLISTER PACK		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076134	06/23/2009	

Labeler - Ohm Laboratories Inc. (184769029)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations

Ohm Laboratories Inc.		051565745	manufacture(51660-209)