

**LORATADINE- loratadine tablet**  
**Chain Drug Marketing Association 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.

**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**

**QC QUALITY CHOICE®**

**NDC 63868-151-30**

**\* Compare to the active ingredient of Claritin®**

**NON-DROWSY■**

**Allergy Relief**

**Loratadine Tablets USP, 10 mg**

**Antihistamine**

**Indoor & Outdoor Allergies**

**Relief of:**

**Sneezing; Runny Nose**

**Itchy, Watery Eyes**

**Itchy Throat or Nose**

**30 Tablets**

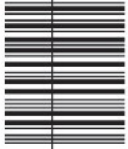
**24 Hour**

**Allergy Relief**

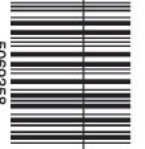
**■ When taken as directed. See Drug Facts Panel.**

**© DISTRIBUTED BY QUALITY CHOICE**

**50693580908**



5069358



5069358

**Drug Facts**

**Active Ingredient (in each tablet)**  
Loratadine USP, 10 mg.....Antihistamine

**Purpose**

**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**  
 ■ 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.

**QC QUALITY CHOICE**

**Allergy Relief**  
**Loratadine Tablets USP, 10 mg**  
**Antihistamine**  
**Indoor & Outdoor Allergies**

**30 Tablets** **24 Hour Allergy Relief**

**QC QUALITY CHOICE**

NDC 63868-151-30

**NON-DROWSY\***

**Allergy Relief**  
**Loratadine Tablets USP, 10 mg**  
**Antihistamine**  
**Indoor & Outdoor Allergies**

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

**30 Tablets** **24 Hour Allergy Relief**

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

\*Compare to the active ingredient of Claritin®

8090

**BUY**  
 10 TABLETS PER BOX  
 5069358

www.schering.com

6 3 5 5 1 5 1 9 5 2 2 5 6

Expiration Date:

Batch No.

**Non Varnish Area**

**Drug Facts (continued)**

**Inactive ingredients** corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

**Questions?** call 1-800-406-7984

See end panel for expiration date.

Keep the carton. It contains important information.

# LORATADINE

loratadine tablet

## Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-151
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)	
STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)	

## 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:63868-151-10	10 in 1 BLISTER PACK		
2	NDC:63868-151-30	30 in 1 BLISTER PACK		
3	NDC:63868-151-01	100 in 1 BOTTLE		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076134	08/19/2003	

**Labeler** - Chain Drug Marketing Association Inc. (011920774)

**Registrant** - Ranbaxy Pharmaceuticals Inc. (937890044)

## Establishment

Name	Address	ID/FEI	Business Operations
------	---------	--------	---------------------

Ohm Laboratories Inc.		051565745	manufacture(63868-151)
-----------------------	--	-----------	------------------------

Revised: 9/2012

Chain Drug Marketing Association Inc.