# LORATADINE ALLERGY RELIEF- loratadine tablet HEB

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**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**

	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° 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** 

## PRINCIPAL DISPLAY PANEL - 10 mg Tablet Blister Pack Carton

*Compare to Claritin*® active ingredient<sup>†</sup>

NDC 37808-529-31

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**Allergy Relief** 

Loratadine Tablets, USP 10 mg Antihistamine

Non-Drowsy\* Allergy Relief

Indoor & Outdoor Allergies

24

Hour

Relief of:

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

## **30 TABLETS**

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

actual

size



Non-Drowsy\* Allergy Relief

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

#### Questions? call 1-800-406-7084 stearate, pregelatinized starch corn starch, lactose monohydrate, magnesium

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Drug Facts (continued)

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- itching of the nose or throat Buizeeus ■
  - IICUY, Watery eyes
  - esou √uunı ■
- or other respiratory anergies.
- temporarily relieves these symptoms due to hay fever

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

Drug Facts



H-E-B

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Non Varnish Area

# **Allergy Relief**

Loratadine Tablets, USP 10 mg **Antihistamine** 

Non-Drowsy\* Allergy Relief

Compare to Claritin® active ingredient<sup>†</sup>



NDC 37808-529-31



# **Allergy Relief**

Loratadine Tablets, USP 10 mg **Antihistamine** 

Non-Drowsy\* Allergy Relief

Indoor & Outdoor Allergies

Relief of:

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

30 TABLETS

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









## LORATADINE ALLERGY RELIEF

loratadine tablet

#### **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-529
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Route of Administration ORAL

## **Active Ingredient/Active Moiety**

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ı	Ingredient Name	Basis of Strength	Strength
ı	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	6 mm	
Flavor		Imprint Code	RX526	
Contains				

F	Packaging				
#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date	
1	NDC:37808-529-11	1 in 1 CARTON	02/01/2018		
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product			
2	NDC:37808-529-31	30 in 1 CARTON	02/01/2018		
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product			
3	NDC:37808-529-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2018		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076134	0 2/0 1/20 18	

# **Labeler** - HEB (007924756)

# Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment			
Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	MANUFACTURE(37808-529)

Revised: 5/2018 HEB