

# LORATADINE ALLERGY RELIEF- loratadine tablet Bryant Ranch Prepack

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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**

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

### **HOW SUPPLIED**

Loratadine 10 mg Tablet

- NDC 63629-7772-1: 20 Tablets in a BOTTLE
- NDC 63629-7772-2: 30 Tablets in a BOTTLE
- NDC 63629-7772-3: 60 Tablets in a BOTTLE
- NDC 63629-7772-4: 14 Tablets in a BOTTLE
- NDC 63629-7772-5: 10 Tablets in a BOTTLE
- NDC 63629-7772-6: 90 Tablets in a BOTTLE
- NDC 63629-7772-7: 28 Tablets in a BOTTLE
- NDC 63629-7772-8: 15 Tablets in a BOTTLE
- NDC 63629-7772-9: 100 Tablets in a BOTTLE

Repackaged/Relabeled by:  
Bryant Ranch Prepack, Inc.  
Burbank, CA 91504

**Loratadine 10 mg Tablet**



GTIN 00363629777211  
 Lot 208620  
 Exp 5/14/2026  
 SN 0123456789

Each tablet contains: Loratadine, USP 10 mg.

Keep this and all drugs out of the reach of children.

Store at 20° to 25°C (68° to 77°F) [See USP Controlled Room Temperature].

Protect from light.

**NDC 63629-7772-1**

**Loratadine Tablets, USP**

**10 mg**

**20 Tablets**



Repackaged by:  
 Bryant Ranch Prepack, Inc.  
 Burbank, CA 91504 USA

Manufactured by:  
 OHM  
 LABORATORIES INC.



## LORATADINE ALLERGY RELIEF

loratadine tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:63629-7772(NDC:51660-526)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>LORATADINE</b> (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	

### Product Characteristics

<b>Color</b>	white (White to Off White)	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	6mm
<b>Flavor</b>		<b>Imprint Code</b>	RX526
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63629-7772-1	20 in 1 BOTTLE; Type 0: Not a Combination Product	07/30/2018	
2	NDC:63629-7772-2	30 in 1 BOTTLE; Type 0: Not a Combination Product	07/02/2018	

3	NDC:63629-7772-3	60 in 1 BOTTLE; Type 0: Not a Combination Product	11/22/2019	
4	NDC:63629-7772-4	14 in 1 BOTTLE; Type 0: Not a Combination Product	12/11/2018	
5	NDC:63629-7772-5	10 in 1 BOTTLE; Type 0: Not a Combination Product	07/05/2018	
6	NDC:63629-7772-6	90 in 1 BOTTLE; Type 0: Not a Combination Product	07/05/2018	
7	NDC:63629-7772-7	28 in 1 BOTTLE; Type 0: Not a Combination Product	04/03/2024	
8	NDC:63629-7772-8	15 in 1 BOTTLE; Type 0: Not a Combination Product	04/03/2024	
9	NDC:63629-7772-9	100 in 1 BOTTLE; Type 0: Not a Combination Product	06/21/2019	

## Marketing Information

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

**Labeler** - Bryant Ranch Prepack (171714327)

**Registrant** - Bryant Ranch Prepack (171714327)

## Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(63629-7772) , RELABEL(63629-7772)

Revised: 4/2024

Bryant Ranch Prepack