

LORATADINE ALLERGY RELIEF- loratadine tablet

Bryant Ranch Prepack

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

NDC: 71335-0100-1: 20 Tablets in a BOTTLE

NDC: 71335-0100-2: 30 Tablets in a BOTTLE

NDC: 71335-0100-3: 60 Tablets in a BOTTLE

NDC: 71335-0100-4: 14 Tablets in a BOTTLE

NDC: 71335-0100-5: 10 Tablets in a BOTTLE

NDC: 71335-0100-6: 90 Tablets in a BOTTLE

NDC: 71335-0100-7: 28 Tablets in a BOTTLE

NDC: 71335-0100-8: 15 Tablets in a BOTTLE

NDC: 71335-0100-9: 100 Tablets in a BOTTLE

Repackaged/Relabeled by:

Bryant Ranch Prepack, Inc.

Burbank, CA 91504

Loratadine 10mg Tablet



GTIN 00371335010010
 Lot 208620
 Exp 5/17/2026
 SN 0123456789

Each tablet contains: Lorradine, USP 10 mg.

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

Store at 20° to 25°C (68° to 77°F); excursions permitted to 15° to 30°C (59° to 86°F) [See USP Controlled Room Temperature].

NDC 71335-0100-1

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

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71335-0100(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:71335-0100-1	20 in 1 BOTTLE; Type 0: Not a Combination Product	02/12/2018	
2	NDC:71335-0100-2	30 in 1 BOTTLE; Type 0: Not a Combination Product	02/23/2018	

3	NDC:71335-0100-3	60 in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2021	
4	NDC:71335-0100-4	14 in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2021	
5	NDC:71335-0100-5	10 in 1 BOTTLE; Type 0: Not a Combination Product	02/20/2018	
6	NDC:71335-0100-6	90 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2018	
7	NDC:71335-0100-7	28 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	
8	NDC:71335-0100-8	15 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	
9	NDC:71335-0100-9	100 in 1 BOTTLE; Type 0: Not a Combination Product	04/05/2024	

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(71335-0100) , RELABEL(71335-0100)

Revised: 4/2024

Bryant Ranch Prepack