

**ALLERGY RELIEF NON DROWSY- loratadine tablet
Bryant Ranch Prepack**

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
- sneezing
- itchy, watery eyes
- 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 (1-800-222-1222) 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 at 20°-25°C (68°-77°F) (see USP Controlled Room Temperature)
- protect from light

Inactive ingredients

lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

Questions or comments?

Call **1-888-588-1418** Monday-Friday 9AM-5PM EST

HOW SUPPLIED

Loratadine 10 mg Tablet

- NDC 71335-1651-1: 20 Tablets in a BOTTLE
- NDC 71335-1651-2: 30 Tablets in a BOTTLE
- NDC 71335-1651-3: 60 Tablets in a BOTTLE
- NDC 71335-1651-4: 14 Tablets in a BOTTLE
- NDC 71335-1651-5: 10 Tablets in a BOTTLE
- NDC 71335-1651-6: 90 Tablets in a BOTTLE
- NDC 71335-1651-7: 28 Tablets in a BOTTLE
- NDC 71335-1651-8: 15 Tablets in a BOTTLE
- NDC 71335-1651-9: 100 Tablets in a BOTTLE

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

Loratadine 10mg Tablet



GTIN 00371335165116
 Lot 208820
 Exp 1/31/2027
 SN 0123456789

Drug Facts	
Active ingredient (in each tablet) Loratadine 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 (1-800-222-1222) right away.	
Other Information •store at 20°-25°C (68°-77°F) (see USP Controlled Room Temperature) •protect from light	
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.	
Inactive Ingredients lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate.	

NDC 71335-1651-1

Loratadine Tablets, USP

10 mg

20 Tablets



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

Manufactured by:
Camber Consumer Care



Package Insert

ALLERGY RELIEF NON DROWSY

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71335-1651(NDC:69230-317)
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
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	439
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71335-1651-1	20 in 1 BOTTLE; Type 0: Not a Combination Product	07/27/2020	
2	NDC:71335-1651-2	30 in 1 BOTTLE; Type 0: Not a Combination Product	07/16/2020	
3	NDC:71335-	60 in 1 BOTTLE; Type 0: Not a Combination	10/16/2020	

3	1651-3	Product	10/10/2020	
4	NDC:71335-1651-4	14 in 1 BOTTLE; Type 0: Not a Combination Product	05/25/2022	
5	NDC:71335-1651-5	10 in 1 BOTTLE; Type 0: Not a Combination Product	01/12/2021	
6	NDC:71335-1651-6	90 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
7	NDC:71335-1651-7	28 in 1 BOTTLE; Type 0: Not a Combination Product	01/30/2025	
8	NDC:71335-1651-8	15 in 1 BOTTLE; Type 0: Not a Combination Product	01/30/2025	
9	NDC:71335-1651-9	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/14/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075209	12/27/2019	12/27/2025

Labeler - Bryant Ranch Prepack (171714327)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-1651) , RELABEL(71335-1651)

Revised: 1/2025

Bryant Ranch Prepack