

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

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**788S (658)**

**Active ingredient (in each tablet)**

Loratadine, USP 10 mg

**Purpose**

Antihistamine

**Uses**

temporarily relieves these symptoms due to hay fever and other upper respiratory allergies:

- runny nose
- sneezing
- itching of the nose and throat
- itchy, watery eyes

**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 12 years and over	1 tablet daily; not more than 1 tablet in 24 hours
children under 12	

children under 12 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

### Other information

- TAMPER EVIDENT: Do not use if imprinted seal under cap is missing or broken.
- store between 20° to 25°C (68° to 77°F)
- protect from light

### Inactive ingredients

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

### Questions?

call **1-800-540-3765**

### HOW SUPPLIED

Loratadine, USP 10 mg

- NDC 72162-2181-9: 90 Tablets in a BOTTLE

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

### Loratadine 10mg Tablets

 GTN Lot Exp SN	<b>Drug Facts</b>				
	<table border="1"> <thead> <tr> <th>Active ingredient (in each tablet)</th> <th>Purpose</th> </tr> </thead> <tbody> <tr> <td>Loratadine, USP 10 mg</td> <td>Antihistamine</td> </tr> </tbody> </table>	Active ingredient (in each tablet)	Purpose	Loratadine, USP 10 mg	Antihistamine
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	Loratadine, USP 10 mg	Antihistamine			
	<b>Uses</b> temporarily relieves these symptoms due to hay fever and other upper respiratory allergies: •runny nose •sneezing •itching of the nose and throat •itchy, watery eyes.				
	<b>Warnings</b> 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.				
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<b>Inactive Ingredients</b> Lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate.					
 Package Insert					

**NDC 72162-2181-9**

**Allergy Relief  
Loratadine Tablets**

**10 mg**

**90 Tablets**



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

**Manufactured by:**  
GERI-CARE  
PHARMACEUTICAL  
CORP



## ALLERGY RELIEF

loratadine tablet

### Product Information

**Product Type**

HUMAN OTC DRUG

**Item Code (Source)**

NDC:72162-2181(NDC:57896-658)

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
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

### 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:72162-2181-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2020	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075209	02/01/2020	

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

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

### Establishment

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(72162-2181) , RELABEL(72162-2181)