

BERKLEY AND JENSEN ALLERGY RELIEF- loratadine tablet BJWC

BJWC Allergy Relief 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 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 age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- do not use if printed foil under cap is broken or missing
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

Questions or comments?

1-800-934-1204

Principal Display Panel

Compare to the active ingredient in Claritin®

berkley jensen®

ORIGINAL PRESCRIPTION STRENGTH

24 HOUR • NON-DROWSY*

ALLERGY RELIEF

LORATADINE TABLETS, 10 mg

ANTIHISTAMINE

INDOOR & OUTDOOR ALLERGIES

24 HOUR RELIEF OF:

- SNEEZING
- RUNNY NOSE
- ITCHY, WATERY EYES
- ITCHY THROAT OR NOSE

365 TABLETS

ACTUAL SIZE

*When taken as directed. See Drug Facts Panel.

Compare to the active ingredient in Claritin^{®**}



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EU #31446 V4
Mant. #0799008



Die Cut

EFC 1548
612-002 EFC1548-1 REV2
(612)



Drug Facts	
Active ingredient (in each tablet) Loratadine 10 mg.....Arthritis name	
Use: Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: <ul style="list-style-type: none"> ■ sneezing ■ itchy, watery eyes ■ itchy throat or nose 	
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 Poison Control Center right away. (1-800-222-1222)	
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Questions or comments? 1-800-BJS-CLUB	
Contains 1 Bottle with 365 Tablets for 365 Days of Relief Distributed by: BJS Wholesale Club, Inc. 350 Campus Drive Methuen, MA 01752 Every product comes with our 100% MONEY-BACK GUARANTEE QUESTIONS? CustomerCare@bjs.com	

berkley jensen®

NON-DROWSY*

ALLERGY RELIEF

LORATADINE TABLETS, 10 mg

ANTIHISTAMINE

INDOOR & OUTDOOR ALLERGIES

24 HOUR RELIEF OF:

SNEEZING • RUNNY NOSE

ITCHY, WATERY EYES

ITCHY THROAT OR NOSE

365 TABLETS

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BERKLEY AND JENSEN ALLERGY RELIEF

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68391-314
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)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	

Product Characteristics

Color	WHITE	Score	no score
Shape	OVAL	Size	8mm
Flavor		Imprint Code	L612
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68391-314-87	1 in 1 PACKAGE	04/17/2009	08/01/2015
1		300 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:68391-314-88	1 in 1 PACKAGE	09/09/2011	
2		365 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:68391-314-79	2 in 1 PACKAGE	09/04/2025	
3	NDC:68391-314-82	200 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076301	04/17/2009	

Labeler - BJWC (159082692)

Revised: 2/2026

BJWC