ALLERGY RELIEF- loratadine capsule, liquid filled CHAIN DRUG CONSORTIUM

1195A-PRV-2020-0903

Drug Facts

Active Ingredient (in each capsule)

Loratadine 10mg

Purpose

Antihistamine

Uses

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

- runny nose
- sneezing
- itchy, water 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 right away (1-800-222-1222).

Directions

adults and children 6 years and over	1 capsule daily; not more than 1 capsule in 24
	hours
children under 6 years of age	ask a doctor
consumers with liver or kidney	ask a doctor
disease	

Other information

- Store between 20-25°C (68-77°F)
- protect from freezing
- retain carton for complete information and warnings

Inactive ingredients

FD&C Blue no. 1, gelatine, mono anddiglyceride of caprilic/capric acid, pharmaceutical ink, polysorbate 80, povidone, purified water, sorbitol, sorbitan solution.

Questions or comments?

1-844-705-4384

PRINCIPAL DISPLAY PANEL

Premier Value®

COMPARE TO THE ACTIVE INGREDIENT IN CLARITIN® LIQUI-GELS®†

*When taken as directed.

See Drug Facts Panel.

NON-DROWSY*

Allergy Relief

LORATADINE 10 MG • 24 HOUR RELIEF

ANTIHISTAMINE

Relieves:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Throar or Nose

10 SOFTGELS**

(**LIQUID-FILLED CAPSULES)



This product is not manufactured or distributed by Bayer Healthcare LLC, distributor of Claritin® or Catalent Pharma Solutions, Inc., owner of the registered trademark Liqui-Gels®. DO NOT USE IF BLISTER UNITS ARE TORN OR BROKEN

Distributed By:
Pharmacy Value Alliance, LLC
407 East Lancaster Avenue
Wayne, PA 19087
Made in India

If for any reason you are not satisfied this product, please return it to the st where purchased for a full refund.

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Questions or comments?

80, povidone, purified water, sorbital sorbitan solution caprylic/capric acid, pharmaceutical ink, polysorbate FD&C blue #1, gelatin, mono and diglycende of Inactive ingredients

and warnings ■ retain carton for complete product information

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 - Itcny, watery eyes esou Kuuru = or other upper respiratory allergies:

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Loratadine 10 mg.

(əınsdeo yotə uı) Active ingredient

Drug Facts





NON-DROWSY* Allergy Relief

Antihistamine

Purpose

LORATADINE 10 MG • 24 HOUR RELIEF ANTIHISTAMINE



COMPARE TO THE ACTIVE INGREDIENT IN CLARITIN® LIQUI-GELS®†

> *When taken as directed. See Drug Facts Panel.

NON-DROWSY* Allergy Relief

LORATADINE 10 MG • 24 HOUR RELIEF ANTIHISTAMINE

Relieves:

- Sneezing
- Itchy, Watery Eyes
- Runny Nose
- Itchy Throat or Nose

Indoor & Outdoor **Allergies**



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ALLERGY RELIEF

loratadine capsule, liquid filled

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:68016-055

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name
Basis of Strength
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)
LORATADINE
10 mg

Inactive Ingredients

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Ingredient Name	Strength	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
GELATIN (UNII: 2G86QN327L)		
CAPRIC ACID (UNII: 4G9EDB6V73)		
POVIDONE K30 (UNII: U725QWY32X)		
WATER (UNII: 059QF0KO0R)		
SORBITOL (UNII: 506T60A25R)		
POLYSORBATE 80 (UNII: 6OZP39ZG8H)		
SORBITAN (UNII: 6092ICV9RU)		

Product Characteristics

Color	blue (Light Blue)	Score	no score
Shape	OVAL (oval shaped)	Size	3mm
Flavor		Imprint Code	21
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016- 055-10	1 in 1 CARTON	12/01/2018	07/31/2025
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA206214	12/01/2018	07/31/2025	

Labeler - CHAIN DRUG CONSORTIUM (101668460)

Revised: 10/2024 CHAIN DRUG CONSORTIUM