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 disease	ask a doctor

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

Indoor Outdoor Allergies

10 SOFTGELS**

(**LIQUID-FILLED CAPSULES)

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**	**Strength**
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients	
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: 6O92ICV9RU)	

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