

CLARITIN- loratadine tablet
Bayer HealthCare LLC.

Clairtin - Project Fortify

Drug Facts

Active ingredient (in each tablet)

Loratadine, 10 mg USP

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

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

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

Stop use

Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children

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

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)

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

Questions or comments?

1-800-CLARITIN (1-800-252-7484) or www.claritin.com

Carton label 10 count tablets

Original Prescription Strength

Non-Drowsy*

Claritin®

loratadine tablets 10 mg/antihistamine

Indoor & Outdoor

Allergies

24

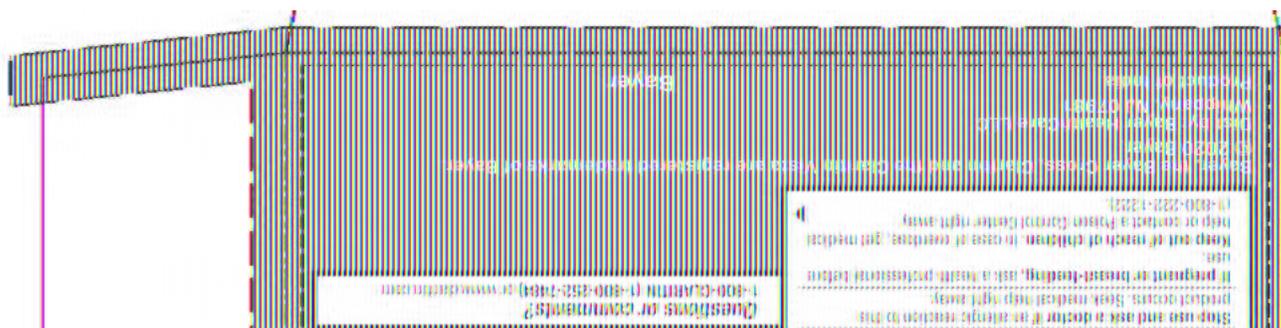
Hour

Relief of:

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

**When taken as directed.*

See Drug Facts Panel





CLARITIN

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11523-0800	
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)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)				
Product Characteristics				
Color	white	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	Claritin;10;458	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-0800-1	1 in 1 CARTON	03/16/2022	
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11523-0800-2	100 in 1 BOTTLE; Type 0: Not a Combination Product	03/16/2022	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA075209	12/01/2020		

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 12/2025

Bayer HealthCare LLC.