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 20º -25º C (68º -77º 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

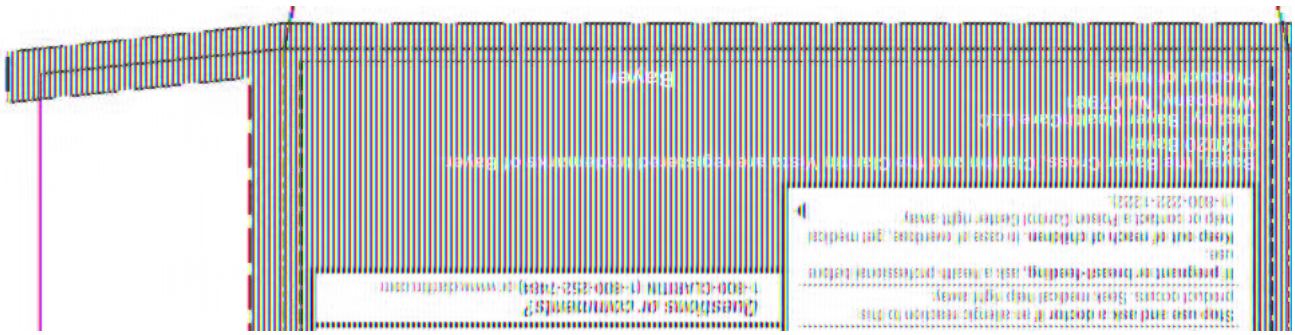
Hour

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: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	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: 3/2023 Bayer HealthCare LLC.