CLARITIN- loratadine tablet tablet Lil' Drug Store Products, Inc.

Claritin ®

Drug Facts

Drug Facts

Active Ingredient

Active Ingredient (in each tablet)

Loratadine, USP 10 mg

Purpose

Purpose

Antihistamine

Uses

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

Do not use if you have ever had an allergic reaction to this product or any of its ingredients.

Ask a doctor before use

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

■ 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

Original Prescription Strength

Non-Drowsy*®

Claritin ®

loratadine tablets 10 mg/antihistamine

24

Hour

Relief of:

- Sneezing
- Runny Nose
- Itchy,

Watery Eyes

• Itchy Throat

or nose

Indoor & Outdoor

Allergies

[Bayer cross] *When takenas directed.

See Drug Facts Panel.

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Tablet [tablet image] [Lil' Drug Store logo]



CLARITIN

loratadine tablet tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66715-9742

Route of Administration ORAL

Active Ingredient/Active Moiety

CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)

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

Inactive Ingredients				
Ingredient Name	Strength			
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				

Product Characteristics				
Color	white	Score	no score	
Shape	ROUND	Size	10mm	
Flavor		Imprint Code	CLARITIN10;458	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:66715- 9742-1	1 in 1 CARTON	06/01/2022		
1		1 in 1 POUCH; Type 0: Not a Combination Product			
	NDC.6671E				

2	NDC:00/13- 9742-2	2 in 1 CARTON	06/01/2022	
2		1 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075209	06/01/2022	

Labeler - Lil' Drug Store Products, Inc. (093103646)

Revised: 12/2022 Lil' Drug Store Products, Inc.