

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 taken as 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

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg

Inactive Ingredients

Ingredient Name	Strength
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
MAGNESIUM STEARATE (UNII: 70097M6130)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

Product Characteristics

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

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:66715			

2	NDC: 66715-9742-2	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: 12/2024

Lil' Drug Store Products, Inc.