LORATADINE - loratadine tablet Cardinal Health

Loratadine Tablets USP 10 mg

Drug Facts

Active ingredient (in each tablet)

Loratadine USP 10 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- itchy, watery eyes
- sneezing
- 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 (1-800-222-1222) right away.

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	ask a doctor
disease	ask a doctor

Other information

- Safety sealed: do not use if the individual blister unit is open or torn
- store at 20° to 25°C (68° to 77°F)
- protect from excessive moisture

Inactive ingredients

lactose monohydrate, magnesium stearate, pregelatinized starch (maize), sodium starch glycolate.

Questions or comments?

call **1-855-274-4122**

DISTRIBUTED BY CARDINAL HEALTH DUBLIN, OHIO 43017 www.myleader.com 1-800-200-6313

Made in India

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 10 mg Blister Carton (20 Tablets)

LEADER™ NDC 70000-0583-1 Non-Drowsy* | 24 Hour Allergy Relief Loratadine Tablets USP, 10 mg | Antihistamine Indoor & Outdoor Allergies

Relief of:

Sneezing; Runny Nose; Itchy, Watery Eyes; Itchy Throat or Nose

COMPARE TO CLARITIN® active ingredient** 100% Money Back Gaurantee

20 TABLETS

Actual Size

*When taken as directed. See Drug Facts Panel.





LORATADINE

loratadine tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:70000-0583

Route of Administration ORAL

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
STARCH, CORN (UNII: O8232NY3SJ)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			

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1				
	Color	WHITE (White to Off-white)	Score	no score
	Shape	ROUND	Size	6mm

Flavor	Imprint Code	39;L
Contains		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000- 0583-1	2 in 1 CARTON	08/30/2021	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:70000- 0583-2	1 in 1 CARTON	06/14/2024	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category			Marketing End Date
ANDA	ANDA208314	08/30/2021	

Labeler - Cardinal Health (063997360)

Registrant - Aurohealth LLC (078728447)

Establishment			
Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(70000-0583), MANUFACTURE(70000-0583)

Revised: 6/2024 Cardinal Health