

**LORATADINE- loratadine tablet**  
**Preferred Pharmaceuticals Inc.**

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**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 disease	ask a doctor

**Other information**

- Tamper-evident: do not use if foil seal under cap, printed with “SEALED for YOUR PROTECTION” is missing, open or broken
- 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: **AUROHEALTH LLC**  
2572 Brunswick Pike  
Lawrenceville, NJ 08648  
Made in India

**Repackaged By: Preferred Pharmaceuticals Inc.**

Code: TS/DRUGS/22/2009

**PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 10 mg**

**NDC 68788-4103**  
**Non-Drowsy\***  
**Loratadine**  
**Tablets USP 10 mg**  
**Antihistamine**

**24 Hour Relief of:**

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

**Indoor & Outdoor Allergies**

**\*When taken as directed. See Drug Facts Panel.**

Preferred Pharmaceuticals Inc.


**Loratadine Tablets 10mg**  
Generic for Claritin

Active ingredient (in each tablet) Loratadine 10mg.....Antihistamine


**Pkg Size:** Exp Date: ##/##/####  
Lot#: Batch#:

Ins:  
Mfg: Aurobindo Pharma Limited  
Prod#:

**Warning**  
Store at 20°- 25°C (68°- 77°F). See USP Controlled Room Temperature. 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. 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. Keep this and all medication out of the reach of children. If pregnant or breast feeding, ask a health professional before use. Tablet is round, white, imprinted with 39 L




CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed.



Directions English

Use as directed by your doctor  
Take \_\_\_ Tablet(s) \_\_\_ time(s) a day



GTIN  
#####  
SN #####  
EXP #####

**Instrucciones Espanol:**  
Usó según lo dirigido por su doctor  
Cada \_\_\_ Tableta(s) \_\_\_ vece(s) al día

Loratadine Tablets 10mg  
Qty: Ins:  
Lot: Bat:  
Prod# (NDC):

Loratadine Tablets 10mg  
Qty: Ins:  
Lot: Bat:  
Prod# (NDC):

Loratadine Tablets 10mg  
Qty:  
Insurance NDC:  
Lot: Bat:

Loratadine Tablets 10mg  
Qty: Ins:  
Lot: Bat:  
Prod# (NDC):

Log
Chart
Billing
Patient

<b>LORATADINE</b>			
loratadine tablet			
Product Information			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68788-4103(NDC:58602-702)
<b>Route of Administration</b>	ORAL		
Active Ingredient/Active Moiety			
<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>	
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg	
Inactive Ingredients			
<b>Ingredient Name</b>	<b>Strength</b>		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
STARCH, CORN (UNII: O8232NY3SJ)			
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)			

## Product Characteristics

<b>Color</b>	WHITE (White to Off-white)	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	6mm
<b>Flavor</b>		<b>Imprint Code</b>	39;L
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-4103-0	10 in 1 BOTTLE; Type 0: Not a Combination Product	04/21/2026	
2	NDC:68788-4103-1	14 in 1 BOTTLE; Type 0: Not a Combination Product	04/21/2026	
3	NDC:68788-4103-5	15 in 1 BOTTLE; Type 0: Not a Combination Product	04/21/2026	
4	NDC:68788-4103-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	04/21/2026	
5	NDC:68788-4103-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/21/2026	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA208314	04/21/2026	

**Labeler** - Preferred Pharmaceuticals Inc. (791119022)

**Registrant** - Preferred Pharmaceuticals Inc. (791119022)

## Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-4103)

Revised: 4/2026

Preferred Pharmaceuticals Inc.