

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

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**Loratadine Tablets, 10 mg**

**ACTIVE INGREDIENT(S)**

Loratadine 10 mg

**PURPOSE**

Antihistamine

**USE(S)**

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

- ☐ runny nose
- ☐ sneezing
- ☐ itchy, water 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 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 DOCTOR IF**

an allergic reaction to this product occurs. Seek medical help right away.

## **PREGNANCY/BREASTFEEDING**

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.

## **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

## **Blister Foil Units**

safety sealed: do not use if the individual blister unit is open or torn

## **STORAGE**

store between 20° to 25°C (68° to 77°F)

## **INACTIVE INGREDIENTS**

Lactose monohydrate, magnesium stearate, sodium starch glycolate and starch maize pregelatinized

## **QUESTIONS OR COMMENTS**

Contact 1-877-770-3183 Mon-Fri 9:00 AM to 4:30 PM EST.

**Repackaged By: Preferred Pharmaceuticals Inc.**

## **PRINCIPAL DISPLAY PANEL**

**Loratadine  
Tablets 10mg**

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

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

Lot#:                      Batch#

Ins:

Mfg: Granules Pharmaceuticals Inc.

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 G 10



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

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



### Directions English

Use as directed by your  
doctor

Take \_\_\_\_\_ tablet(s)  
every \_\_\_\_\_ hours.



GTIN

#####

##### NS

EXP ###/###/###

## Instrucciones Español:

## Uso según lo dirigido por su doctor

• Toma \_\_\_\_\_ tableta(s) \_\_\_\_\_ horas, cada

# LORATADINE

loratadine tablet

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68788-8659(NDC:70010-162)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>LORATADINE</b> (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg

## Inactive Ingredients

Ingredient Name	Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STARCH, CORN (UNII: O8232NY3SJ)	
MAGNESIUM STEARATE (UNII: 70097M6I3O)	
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>	G;10
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8659-0	10 in 1 BOTTLE; Type 0: Not a Combination Product	05/13/2024	

2	NDC:68788-8659-1	14 in 1 BOTTLE; Type 0: Not a Combination Product	05/13/2024	
3	NDC:68788-8659-5	15 in 1 BOTTLE; Type 0: Not a Combination Product	05/13/2024	
4	NDC:68788-8659-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	05/13/2024	
5	NDC:68788-8659-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	05/13/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210722	05/13/2024	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-8659)