#### LORATADINE ANTIHISTAMINE- loratadine tablet Preferred Pharmaceuticals Inc.

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## Perrigo Loratadine Tablets, 10 mg Drug Facts

#### Active ingredient (in each tablet)

Loratadine 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 right away. (1-800-222-1222)

# Directions

adults and children 6 years and	1 tablet daily; not more than 1 tablet in 24 hours
over	
children under 6 years of age	ask a doctor
consumers with liver or kidney	ask a doctor
disease	

## Other information

- do not use if printed foil under cap is broken or missing
- store between 20° to 25°C (68° to 77°F)

## **Inactive ingredients**

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

## Questions or comments?

1-800-719-9260

## **Principal Display Panel**

Compare to Claritin<sup>®</sup> active ingredient

Original Prescription Strength

Non-Drowsy\*

Loratadine Tablets, 10 mg

Antihistamine

Indoor & Outdoor Allergies

24 Hour Relief of:

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

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

actual size

Padagis

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Loratadine Tablets 10mg	PREFERRED Pharmaceuticals, Inc	CAUTION: Federal law PROHIBITS transfer of this drug to any person other thean the patient for whom it was prescribed	Loratadine Tablets 10mg Qty: Ins: Lot#: Bat#: Prod# (NDC):	Log
Generic for Claritin Active ingredient (in each tablet) Loratadine 10mgAntihistamine <b>Pkg Size:</b> Exp Date: Lot#: Batch#:	 your )		Loratadine Tablets 10mg Qty: Ins: Lot#: Bat#: Prod# (NDC):	Chart
Ins: Mfg: Padagis Israel Pharmaceuticals Ltd Prod#: W	Directions English Directed by yo tablet(s) Directed by yo	Instrucciones Espandod doctor tableta(s) doctor tableta(s)horas.	Loratadine Tablets 10mg Qty: Insurance NDC: Lot#: Bat#:	Billing
Store at 20°-25°C (6°-Warning Room et al. 20°-25°C (6°-Warning allergic reaction to this product or any of its ingredients Ask a doctor before use if you have itver or kidney disease. When using this product do not take more than drowsiness. Store use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right way face this and all medication out of the reach of professional before use. Tablet is oval, white, imprinted with Lo12	Use as dirr Use as dirr doctor Take every	Instru Uso según por su doc Toma cada	Loratadine Tablets 10mg Qty: Ins: Lot#: Bat#: Prod# (NDC):	Patient

LORATADINE AN loratadine tablet	ITIHISTAMI	NE						
Product Information	n							
Product Type	HUMAN OTC DRUG Item Code (Source) NDC:68788-8					8-8629(NDC	-8629(NDC:45802-650)	
Route of Administratio	n ORAL							
Active Ingredient/Ac	tive Moiety							
Ingredient Name Basis of Str						trength	Strength	
LORATADINE (UNII: 7AJO3B	07QN) (LORATADINE	E - UNII:7	AJO3BO7QN)		LORATADINE		10 mg	
Inactive Ingredients	<b>Ingredie</b> (UNII: EWQ57Q8I5X NII: 70097M6I30)	()	16			St	rength	
Product Characteris	itics							
Color	WHITE Score no score							
Shape	OVAL					8mm	3mm	
					L612			
Contains								
Packaging								
# Item Code	Package Des	scriptio	on		eting Start Date		eting End Date	

	Category	Citation		Dutt		
	Marketing	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
Marketing Information						
5	NDC:68788- 8629-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2024			
4	NDC:68788- 8629-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2024			
3	NDC:68788- 8629-5	15 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2024			
2	NDC:68788- 8629-1	14 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2024			
1	NDC:68788- 8629-0	10 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2024			

Labeler - Preferred Pharmaceuticals Inc. (791119022)

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

# Establishment

Name	Address	ID/FEI	<b>Business Operations</b>
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-8629)

Revised: 4/2024

Preferred Pharmaceuticals Inc.