

ALLERGY RELIEF NON DROWSY- loratadine tablet
Preferred Pharmaceuticals, Inc.

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
- sneezing
- itchy, watery 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 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

- store at 20°-25°C (68°-77°F) (see USP Controlled Room Temperature)
- protect from light

Inactive ingredients

lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

Questions or comments?

Call **1-888-588-1418** Monday-Friday 9AM-5PM EST

Principal Display Panel

†Compare to the active ingredient in Claritin® 24 Hour

Non-drowsy*

Allergy Relief

Loratadine Tablets, USP 10 mg / Antihistamine

Indoor & outdoor allergies

24 Hour Relief of:

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

Tablets

Gluten-Free

***When taken as directed. See Drug Facts panel.**

†This product is not manufactured or distributed by Bayer Healthcare LLC, distributor of Claritin® 24 Hour

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAPS IS BROKEN OR MISSING.

Camber Consumer Care Inc., Piscataway, NJ 08854, USA

Repackaged By: Preferred Pharmaceuticals Inc.

Package Label

Loratadine Tablets 10mg

Generic for: Claritin

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

Pkg Size: Exp Date:

Lot#:

Batch#:

Ins:

Mfg: Camber Consumer Care

Prod#:

Warning

Store at 20°-25°C (68°-77°F). See USP Controlled Room Temperature. Protect from light. 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 439



Directions English

Take _____ tablet(s)
every _____ hours.

Use as directed by your
doctor



Instrucciones Español:

Toma _____ tableta(s)
cada _____ horas.

Use según lo dirigido
por su doctor

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: Ins:

Insurance NDC:

Lot#: Bat#:

Loratadine Tablets 10mg

Qty: Ins:

Lot#: Bat#:

Prod# (NDC):

CAMBER CONSUMER CARE Non-Drowsy Allergy Relief

ALLERGY RELIEF NON DROWSY

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68788-7769(NDC:69230-317)
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
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)			
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)			
Product Characteristics			
Color	WHITE	Score	no score

Shape		ROUND	Size		6mm
Flavor			Imprint Code		439
Contains					
Packaging					
#	Item Code	Package Description		Marketing Start Date	Marketing End Date
1	NDC:68788-7769-0	10 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		08/03/2020	
2	NDC:68788-7769-1	14 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		08/03/2020	
3	NDC:68788-7769-5	15 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		08/03/2020	
4	NDC:68788-7769-3	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		08/03/2020	
5	NDC:68788-7769-9	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		08/03/2020	
Marketing Information					
Marketing Category		Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
ANDA		ANDA075209		08/03/2020	

Labeler - Preferred Pharmaceuticals, Inc. (791119022)

Registrant - Preferred Pharmaceuticals, Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals, Inc.		791119022	REPACK(68788-7769)

Revised: 7/2023

Preferred Pharmaceuticals, Inc.