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 UPS 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.

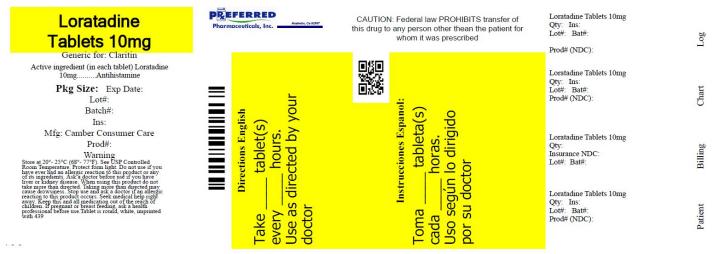
 $\dagger This product is not manufactured or distributed by Bayer Healthcare LLC, distributor of Claritin <math display="inline">\ensuremath{\mathbb{R}}$ 24 Hour

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



CAMBER CONSUMER CARE Non-Drowsy Allergy Relief

ALLERGY RELIEF N loratadine tablet	ION DROWS	Y					
Product Information							
Product Type	HUMAN OTC DRU	G Item Code (Sourc	e) NDC:6878	8-7769(ND0	2:69230-317)		
Route of Administration	ORAL						
Active Ingredient/Activ	e Moiety						
Ingredient Name Basis of Strengt				trength	Strength		
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN) LORATADINE					10 mg		
Inactive Ingredients							
Ingredient Name					Strength		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)							
MAGNESIUM STEARATE (UNII:	MAGNESIUM STEARATE (UNII: 70097M6I30)						
MICROCRYSTALLINE CELLULO	DSE (UNII: OP1R32D6	1U)					
SODIUM STARCH GLYCOLATE	TYPE A CORN (UNII	: AG9B65PV6B)					
Product Characteristic	S						
Color	VHITE S	core		no score			

Shape ROUND		ROUND	Size	6mm		
Flavor			Imprint Code		439	
Со	ntains					
Pa	ackaging					
#	ltem Code		Package Description		Marketing Start Date	Marketing End Date
	NDC:68788- 7769-0	10 in 1 BO Combinatio	TTLE, PLASTIC; on Product	Type 0: Not a	08/03/2020	
	NDC:68788- 7769-1	14 in 1 BO Combinatio	TTLE, PLASTIC; on Product	Type 0: Not a	08/03/2020	
	NDC:68788- 7769-5	15 in 1 BO Combinatio	TTLE, PLASTIC; on Product	Type 0: Not a	08/03/2020	
	NDC:68788- 7769-3	30 in 1 BO Combinatio	TTLE, PLASTIC; on Product	Type 0: Not a	08/03/2020	
	NDC:68788- 7769-9	90 in 1 BO Combinatio	TTLE, PLASTIC; on Product	Type 0: Not a	08/03/2020	
М	arketing	Inform	nation			
	Marketing Category		lication Num	ber or Monograph ation	Marketing Start Date	Marketing End Date
AN	DA	ANDA0	75209		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.