ALLERGY RELIEF- loratadine tablet
Preferred Pharmaceuticals Inc.

788S (658)

Active ingredient (in each tablet)

Loratadine, USP 10 mg

Purpose

Antihistamine

Uses

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

- runny nose
- sneezing
- itching of the nose and throat
- itchy, watery eyes

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.

Directions

adults and children 12 years and over	1 tablet daily; not more than 1 tablet in 24 hours

children under 12 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- store between 20° to 25°C (68° to 77°F)
- protect from light

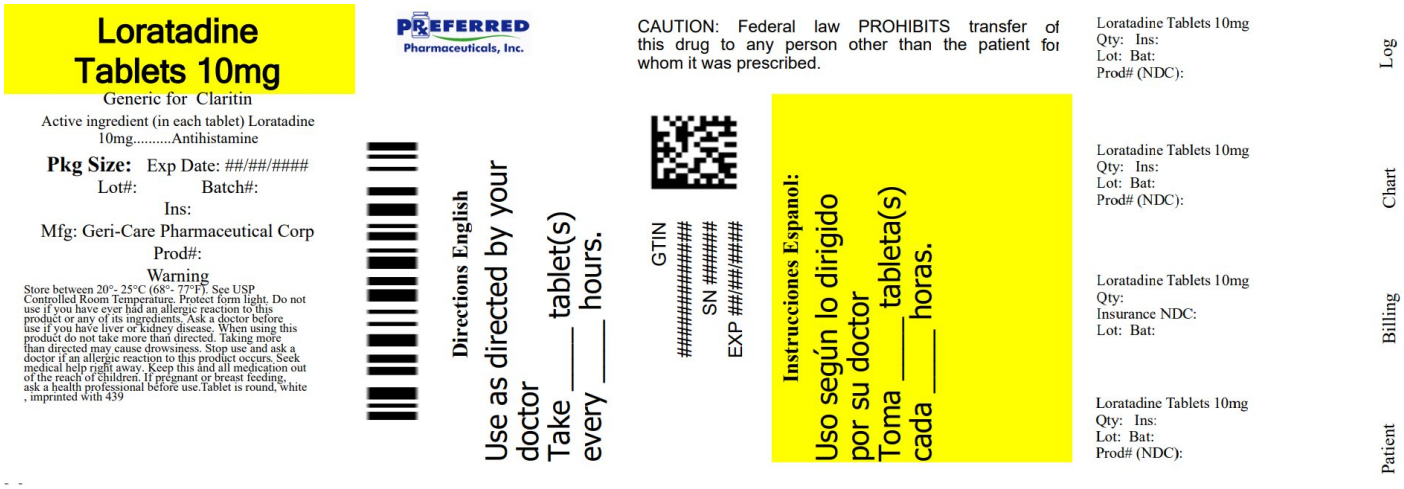
Inactive ingredients

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

Questions?

call **1-800-540-3765**

package label



ALLERGY RELIEF

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68788-4016(NDC:57896-658)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients

Ingredient Name	Strength
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

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:68788-4016-0	10 in 1 BOTTLE; Type 0: Not a Combination Product	08/29/2025	
2	NDC:68788-4016-1	14 in 1 BOTTLE; Type 0: Not a Combination Product	08/29/2025	
3	NDC:68788-4016-5	15 in 1 BOTTLE; Type 0: Not a Combination Product	08/29/2025	
4	NDC:68788-4016-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	08/29/2025	
5	NDC:68788-4016-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	08/29/2025	
6	NDC:68788-4016-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	08/29/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075209	08/29/2025	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-4016)