LORATADINE- loratadine tablet
Preferred Pharmaceuticals Inc.

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 20º to 25ºC (68º to 77ºF)

INACTIVE INGREDIENTS

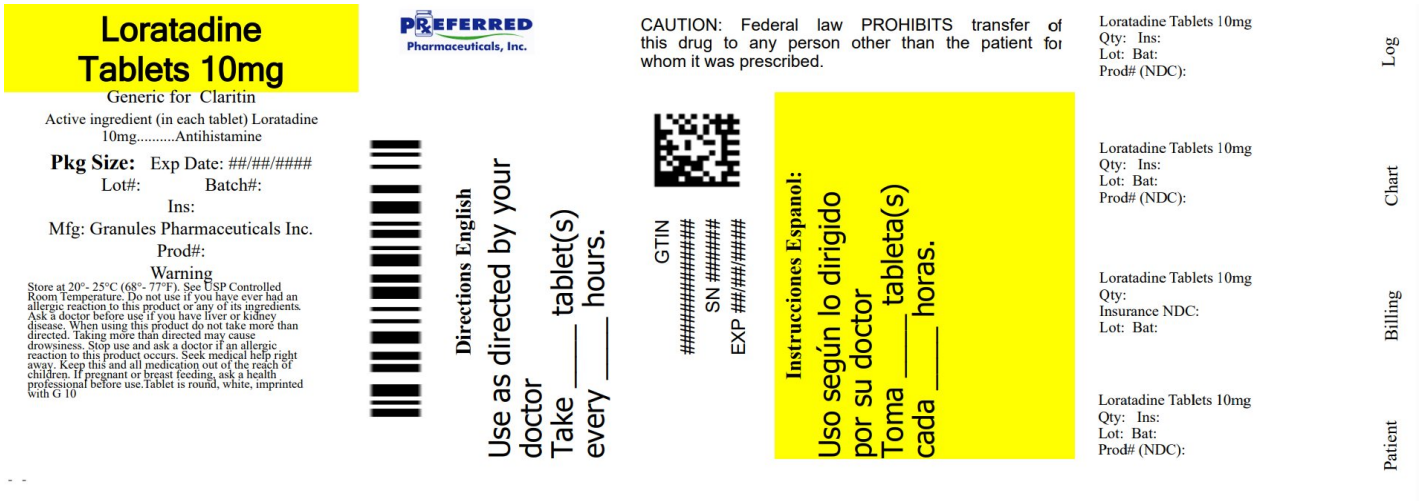
Lactose monohydrate, magnesium stearate, sodium starch glycolate and starch maize pregelatinized

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

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68788-8659(NDC:70010-162)
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
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STARCH, CORN (UNII: O8232NY3SJ)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	white (White to off white)	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	G;10
Contains			

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)

Revised: 7/2025 Preferred Pharmaceuticals Inc.