LORATADINE- loratadine tablet
Preferred Pharmaceuticals Inc.

Loratadine tablets USP, 10mg/antihistamine

ACTIVE INGREDIENT(S)

Active ingredient (in each tablet)
Loratadine USP, 10 mg

PURPOSE

Antihistamine

USE(S)

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.

ASK A DOCTOR OR PHARMACIST BEFORE USE IF

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

Bottles:

- Tamper-evident: do not use if foil seal under cap, printed with "SEALED for YOUR PROTECTION" is missing, open or broken
- store between 20° to 25°C (68° to 77°F)

Blisters packs:

- safety sealed: do not use if the individual blister unit imprinted with loratadine is open or torn
- store between 20° to 25°C (68° to 77°F)
- protect from excessive moisture

INACTIVE INGREDIENTS

corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

Repackaged By: Preferred Pharmaceuticals Inc.

QUESTIONS OR COMMENTS?

Call 1-888-588-1418

PRINCIPAL DISPLAY PANEL

Loratadine-tablets-USP-10mg-30's-carton

LORATADINE

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68788-8605(NDC:69230-323)
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
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	white (White to Off white)	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	H;L20
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8605-0	10 in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024	

2	NDC:68788-8605-1	14 in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024	
3	NDC:68788-8605-5	15 in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024	
4	NDC:68788-8605-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024	
5	NDC:68788-8605-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	03/13/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211718	03/13/2024	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-8605)

Revised: 7/2025 Preferred Pharmaceuticals Inc.