LORATADINE ALLERGY RELIEF- loratadine tablet
A-S Medication Solutions

Loratadine Allergy Relief

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
- 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.

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 (1-800-222-1222).

Directions

adults and children 6 years and over	1 tablet daily; not more than 1 tablet in 24 hours
children under 6 years of	ask a doctor

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 excessive moisture
- **TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL IS BROKEN OR MISSING FROM BOTTLE.**

Inactive ingredients

corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

Manufactured for/ Distributed by:

Marlex Pharmaceuticals, Inc.

New Castle, DE 19720

Rev. 10/22 SP

LORATADINE ALLERGY RELIEF

LORATADINE ALLERGY RELIEF

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50090-7507(NDC:10135-763)
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
ZEA MAYS (CORN) STARCH (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**	RX526
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090-7507-0	10 in 1 BOTTLE; Type 0: Not a Combination Product	02/18/2025	
2	NDC:50090-7507-5	90 in 1 BOTTLE; Type 0: Not a Combination Product	02/18/2025	
3	NDC:50090-7507-4	30 in 1 BOTTLE; Type 0: Not a Combination Product	02/18/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076134	10/01/2022	

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-7507)

Revised: 2/2025 A-S Medication Solutions