ALLERGY RELIEF NON DROWSY- loratadine tablet
A-S Medication Solutions

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 20º-25ºC (68º-77ºF) (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

HOW SUPPLIED

Product: 50090-5067

NDC: 50090-5067-5 90 TABLET in a BOTTLE

NDC: 50090-5067-0 10 TABLET in a BOTTLE

NDC: 50090-5067-1 20 TABLET in a BOTTLE

NDC: 50090-5067-3 15 TABLET in a BOTTLE

NDC: 50090-5067-4 30 TABLET in a BOTTLE

NDC: 50090-5067-6 7 TABLET in a BOTTLE

Loratadine

ALLERGY RELIEF NON DROWSY

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50090-5067(NDC:69230-317)
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)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	

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:50090-5067-3	15 in 1 BOTTLE; Type 0: Not a Combination Product	05/29/2020	

2	NDC:50090-5067-5	90 in 1 BOTTLE; Type 0: Not a Combination Product	05/29/2020	
3	NDC:50090-5067-0	10 in 1 BOTTLE; Type 0: Not a Combination Product	05/29/2020	
4	NDC:50090-5067-4	30 in 1 BOTTLE; Type 0: Not a Combination Product	05/29/2020	
5	NDC:50090-5067-6	7 in 1 BOTTLE; Type 0: Not a Combination Product	05/29/2020	
6	NDC:50090-5067-1	20 in 1 BOTTLE; Type 0: Not a Combination Product	05/29/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075209	12/27/2019	

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-5067) , REPACK(50090-5067)

Revised: 2/2023 A-S Medication Solutions