ALLERGY RELIEF- loratadine tablet
A-S Medication Solutions

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	

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

Other information

- TAMPER EVIDENT: Do not use if imprinted seal under cap is missing or broken.
- 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**

HOW SUPPLIED

Product: 50090-6164

NDC: 50090-6164-4 30 TABLET in a BOTTLE

NDC: 50090-6164-5 90 TABLET in a BOTTLE

NDC: 50090-6164-0 10 TABLET in a BOTTLE

loratadine

ALLERGY RELIEF loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50090-6164(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:50090-6164-0	10 in 1 BOTTLE; Type 0: Not a Combination Product	10/12/2022	
2	NDC:50090-6164-4	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/12/2022	
3	NDC:50090-6164-5	90 in 1 BOTTLE; Type 0: Not a Combination Product	10/12/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075209	02/01/2020	

Labeler - A-S Medication Solutions (830016429)

Establishment

Name	Address	ID/FEI	Business Operations
A-S Medication Solutions		830016429	RELABEL(50090-6164) , REPACK(50090-6164)