ALLERGY RELIEF- loratadine tablet
Allegiant Health

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309 - Health A2Z Allergy Relief

Active ingredient (in each tablet)

Loratadine 10mg

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.

You may report side effects to 1-888-952-0050.

Pregnancy/Breastfeeding

ask a health professional before use.

Keep out of reach of children

In case of accidental overdose, contact a doctor or 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 between 20° to 25°C (68° to 77°F)
- protect from excessive moisture
- do not use if imprinted seal under safety cap is broken or missing

Inactive ingredients

lactose monohydrate, magnesium stearate, sodium starch glycolate
May contain: microcrystalline cellulose, pregelatinized starch

Principal Display Panel

Loratadine

ALLERGY RELIEF

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69168-414
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)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	**MICROCRYSTALLINE CELLULOSE** (UNII: OP1R32D61U)	
May contain	**STARCH, CORN** (UNII: O8232NY3SJ)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69168-414-17	300 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
2	NDC:69168-414-03	3 in 1 CARTON	05/01/2020	
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:69168-414-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	05/18/2022	
4	NDC:69168-414-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020	
5	NDC:69168-414-09	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	05/01/2020	
6	NDC:69168-414-08	5 in 1 BLISTER PACK; Type 0: Not a Combination Product	05/01/2020	
7	NDC:69168-414-02	150 in 1 BOTTLE; Type 0: Not a Combination Product	11/06/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210722	05/01/2020	

Labeler - Allegiant Health (079501930)

Revised: 6/2020 Allegiant Health