ALLERGY RELIEF- loratadine tablet LORATADINE 10MG- loratadine 10mg tablet Allegiant Health

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.

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

*corn starch, lactose monohydrate, magnesium stearate, pregelantinized starch, *sodium starch glycolate

*May contain these ingredients

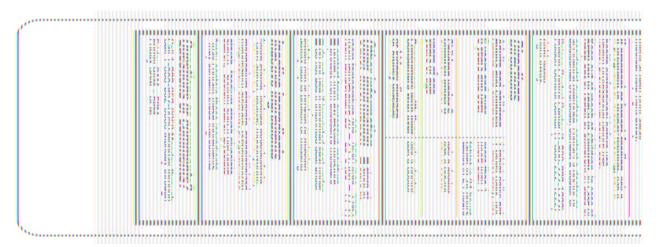
Principal Display Panel



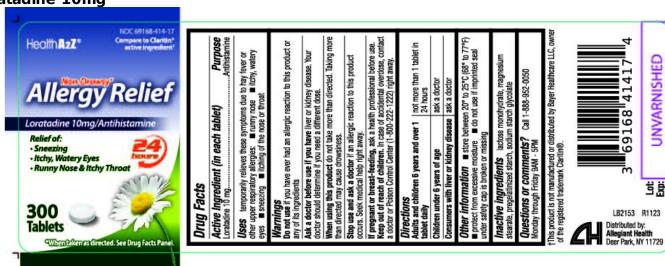


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設置



Loratadine 10mg



Loratadine 10mg

ALLERGY RELIEF loratadine tablet **Product Information HUMAN OTC DRUG Item Code (Source)** NDC:69168-414 **Product Type**

Route of Administration

ORAL

Active Ingredient/Active Moiety

ı	,		
l	Ingredient Name	Basis of Strength	Strength
ı	LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients			
Ingredient Name	Strength		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			

Other Ingredients				
Ingredient Kind Ingredient Name Quantit				
May contain	STARCH, CORN (UNII: O8232NY3SJ)			
May contain	SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)			

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						
NDC:69168- 414-17	300 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020							
NDC:69168- 414-03	3 in 1 CARTON	05/01/2020							
	10 in 1 BLISTER PACK; Type 0: Not a Combination Product								
NDC:69168- 414-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	05/18/2022							
NDC:69168- 414-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2020							
NDC:69168- 414-09	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	05/01/2020							
NDC:69168- 414-08	5 in 1 BLISTER PACK; Type 0: Not a Combination Product	05/01/2020							
NDC:69168- 414-02	150 in 1 BOTTLE; Type 0: Not a Combination Product	11/06/2023							
NDC:69168- 414-99	365 in 1 BOTTLE; Type 0: Not a Combination Product	03/18/2025							
	NDC:69168- 414-17 NDC:69168- 414-03 NDC:69168- 414-60 NDC:69168- 414-30 NDC:69168- 414-09 NDC:69168- 414-08 NDC:69168- 414-02 NDC:69168-	NDC:69168- 414-17 NDC:69168- 414-03 3 in 1 CARTON 10 in 1 BLISTER PACK; Type 0: Not a Combination Product NDC:69168- 414-60 NDC:69168- 414-30 NDC:69168- 414-30 NDC:69168- 414-30 NDC:69168- 414-09 NDC:69168- 414-09 NDC:69168- 414-09 NDC:69168- 414-09 NDC:69168- 414-08 NDC:69168- 414-02 NDC:69168- 365 in 1 BOTTLE; Type 0: Not a Combination Product NDC:69168- 414-02 NDC:69168- 414-02	NDC:69168- 414-03 3 in 1 CARTON 10 in 1 BLISTER PACK; Type 0: Not a Combination Product NDC:69168- 414-60 NDC:69168- 414-30 3 in 1 BOTTLE; Type 0: Not a Combination Product NDC:69168- 414-60 NDC:69168- 414-30 NDC:69168- 414-30 NDC:69168- 414-30 NDC:69168- 414-09 NDC:69168- 414-08 S in 1 BLISTER PACK; Type 0: Not a Combination Product NDC:69168- 414-08 NDC:69168- 414-08 NDC:69168- 414-08 S in 1 BOTTLE; Type 0: Not a Combination Product NDC:69168- 414-08 NDC:69168- 414-02 NDC:69168- 414-02 NDC:69168- 414-02 NDC:69168- 365 in 1 BOTTLE; Type 0: Not a Combination Product NDC:69168- 365 in 1 BOTTLE; Type 0: Not a Combination NDC:69168- 365 in 1 BOTTLE; Type 0: Not a Combination NDC:69168- 365 in 1 BOTTLE; Type 0: Not a Combination NDC:69168-						

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

LORATADINE 10MG

loratadine 10mg tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69168-475

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AIO3BO7ON) (LORATADINE - UNII:7AIO3BO7ON)	LORATADINE	10 mg

Inactive Ingredients

mactive ingredients					
Ingredient Name	Strength				
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)					
MAGNESIUM STEARATE (UNII: 70097M6I30)					

Other Ingredients				
Ingredient Kind	Ingredient Kind Ingredient Name Quantity			
May contain	STARCH, CORN (UNII: O8232NY3SJ)			
May contain	SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)			

Product Characterist	Product Characteristics				
Color	white	Score	no score		
Shape	ROUND	Size	6mm		
Flavor		Imprint Code	H;L20		
Contains					

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69168- 475-02	150 in 1 BOTTLE; Type 0: Not a Combination Product	03/18/2025			
2	NDC:69168- 475-08	1 in 1 CARTON	09/30/2025			
2		5 in 1 BLISTER PACK; Type 0: Not a Combination Product				
2	NDC:69168-	1 in 1 CARTON	00/20/2025			

3	475-09	I III I CARTON	09/30/2023	
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:69168- 475-17	300 in 1 BOTTLE; Type 0: Not a Combination Product	09/30/2025	
5	NDC:69168- 475-99	365 in 1 BOTTLE; Type 0: Not a Combination Product	09/30/2025	
6	NDC:69168- 475-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	09/30/2025	
7	NDC:69168- 475-30	1 in 1 CARTON	09/30/2025	
7		30 in 1 BOTTLE; Type 0: Not a Combination Product		
8	NDC:69168- 475-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	11/11/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211718	03/18/2025	

Labeler - Allegiant Health (079501930)

Revised: 6/2020 Allegiant Health