DG HEALTH ALLERGY RELIEF- loratadine tablet Dolgencorp Inc

Dolgencorp, LLC Allergy Relief Drug Facts

Active ingredient (in each tablet)

Loratadine 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 age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- do not use if blister unit is broken or torn
- store between 20° to 25°C (68° to 77°F)
- protect from excessive moisture

Inactive ingredients

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

Questions or comments?

1-888-309-9030

Principal Display Panel

DG[™]|health

Compare to the active ingredient of Claritin® Tablets

Original Prescription Strength

Allergy Relief

Loratadine Tablets, 10 mg • Antihistamine

Indoor & Outdoor Allergies

24 HOUR

Non Drowsy*

24 Hour Relief of:

- Sneezing Runny nose
- Itchy, watery eyes
- Itchy throat or nose

10 mg

*When taken as directed. See Drug Facts Panel.

30 Tablets

Actual Tablet Size

Original Prescription Strength

lergy Relief

Loratadine Tablets, 10 mg • Antihistamine

30 Tablets for 30 Days of Relief



Compare to the active ingredient of Claritin® Tablets**

Original Prescription Strength

Allergy Relief

Loratadine Tablets, 10 mg • Antihistamine

Indoor & Outdoor Allergies

Non Drowsy*

24 Hour Relief of:

- Sneezing Runny nose
- Itchy, watery eyes
- Itchy throat or nose

10 mg

*When taken as directed. See Drug Facts Panel.



Actual Tablet Size

61239 VT C5



Drug Facts

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- runn y no se itch y, watery eyes
- sneezin g
 - 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.

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Drug Facts (continued)

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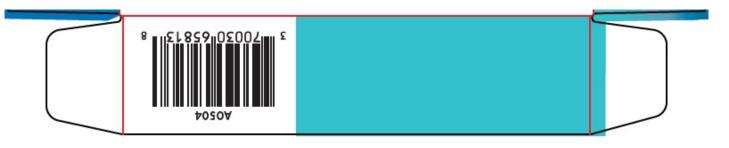
DISTRIBUTED BY OLD EAST MAIN CO. 100 MISSION RIDGE GOODLETTS/ILLE, TN 3707 2

100% Satisfaction Guarante ed! (888)309-9030

**This product is not manufactured or distributed by Bayer HealthCare LLC, distributor of Claritin® Tablets.

Original Prescription Strength oratadine Tablets, 10 mg • Antihistamine

30 Tablets for 30 Days of Relief



DG HEALTH ALLERGY RELIEF

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-806
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg	

Strength

Product Characteristics			
Color WHITE Score no score			
Shape	OVAL	Size	8mm
Flavor		Imprint Code	L612
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:55910-806- 39	30 in 1 CARTON	04/18/2012		
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product			
2	NDC:55910-806- 72	1 in 1 CARTON	04/13/2012		
2		60 in 1 BOTTLE; Type 0: Not a Combination Product			
3	NDC:55910-806- 49	40 in 1 CARTON	04/18/2012		
,		1 in 1 BLISTER PACK; Type 0: Not a Combination			

I	3		Product		
l	4	NDC:55910-806- 65	1 in 1 CARTON	04/28/2023	
l	4		30 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA076301	04/13/2012	

Labeler - Dolgencorp Inc (068331990)

Revised: 10/2023 Dolgencorp Inc