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

Allergy Relief

Loratadine Tablets, 10 mg • **Antihistamine**

30 Tablets for
30 Days of Relief

DG™ | health

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

24 HOUR

10 mg

*When taken as directed.
See Drug Facts Panel.

30 Tablets

Actual Tablet Size

61239 VT C5

CODE AREA

Drug Facts

Active Ingredient (In each tablet) ***Purpose***

Loratadine 10 mg.....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

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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
GOODLETTSVILLE, TN 37072

100% Satisfaction Guaranteed!
(888)309-9030

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

Original Prescription Strength

Allergy Relief

Loratadine 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

Inactive Ingredients	
Ingredient Name	**Strength**
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	

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

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	
3		1 in 1 BLISTER PACK; Type 0: Not a Combination		

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

Marketing Information

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

Labeler - Dolgencorp Inc (068331990)

Revised: 10/2023 Dolgencorp Inc