

LORATADINE ANTIHISTAMINE- loratadine tablet
NuCare Pharmaceuticals, Inc.

Perrigo Loratadine Tablets, 10 mg 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 printed foil under cap is broken or missing
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

Questions or comments?

1-800-719-9260

Principal Display Panel

The image shows the principal display panel for Loratadine 10mg #90 Tablets. The panel is white with a blue header for NuCare Pharmaceuticals, Inc. The product name 'Loratadine 10mg #90 Tablets' is prominently displayed in the center. To the left, there is a vertical barcode and text indicating the package size and manufacturer. To the right, there is a QR code and additional product information. The bottom of the panel contains a warning to keep out of reach of children and storage instructions.

NuCare Pharmaceuticals, Inc.

NDC: 66267-653-90

Loratadine 10mg #90 Tablets

Each tablet contains:
Loratadine 10mg. Antihistamine 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) Oval White Tablet Debossed: "L512" on one side.

Product #: P0653090

WARNING: KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED TEMPERATURE 68-77°F.

Loratadine 10mg
Lot: 000000 NDC: 66267-0653-90
MFR NDC: 45802-650-87 Exp.: 00-00

Loratadine 10mg
Lot: 000000 NDC: 66267-0653-90
MFR NDC: 45802-650-87 Exp.: 00-00

GTIN 00366267653909
Serial# 000000000002
Exp. Date 00-00
LOT#: 000000

Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

Packaged By: NuCare Pharmaceuticals, Inc.
Orange, CA 92867

Patent Instructions:

Take every _____ hours _____ times a day.

66267065390*90*000000*000000

Rev 01/01/19

Distributed by: 3 6 6 2 6 7 6 5 3 9 0
Perrigo Allegan, MI 49010

LORATADINE ANTIHISTAMINE

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66267-653(NDC:45802-650)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	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	L6 12
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66267-653-07	7 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
2	NDC:66267-653-10	10 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
3	NDC:66267-653-14	14 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
4	NDC:66267-653-20	20 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
5	NDC:66267-653-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
6	NDC:66267-653-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
7	NDC:66267-653-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076301	10/15/2008	

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals, Inc.		010632300	repack(66267-653)

Revised: 1/2021

NuCare Pharmaceuticals, Inc.