

**LORATADINE NON DROWSY- loratadine tablet**  
**NuCare Pharmaceuticals, Inc.**

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**Drug Facts**

**Active ingredient (in each tablet)**

Loratadine, USP 10 mg

**Purpose**

Antihistamine

**Uses**

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
- itchy, watery eyes
- 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 (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 at 20°-25°C (68°-77°F) (see USP Controlled Room Temperature)
- protect from light

## Inactive ingredients

lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

## Questions or comments?

Call **1-888-588-1418** Monday-Friday 9AM-5PM EST

## Package Label

 NuCare Pharmaceuticals, Inc.

**NDC: 68071-2208-9**  
**Loratadine 10mg**  
**#90 Tablets**

Each tablet contains: Loratadine, USP 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. In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Round White Tablet Debossed: "439" on one side

Product #: P0653090

Take \_\_\_\_\_ times a day.  
\_\_\_\_\_ every \_\_\_\_\_ hours

Patent Instructions:

Distributed by: 3 68071 22089 6  
Camber Consumer Care Inc.,  
Piscataway, NJ 08854  
Packaged By:  
NuCare Pharmaceuticals, Inc.  
Orange, CA 92867

Rev. 01/01/19

WARNING: KEEP OUT OF REACH OF CHILDREN

Store at controlled temperature 68-77°F.

Loratadine 10mg  
Lot: 00000 NDC: 68071-2208-09  
MFR NDC: 69230-317-03 Exp.: 00-00  
Serial# 0000000002

Loratadine 10mg  
Lot: 00000 NDC: 68071-2208-09  
MFR NDC: 69230-317-03 Exp.: 00-00  
Serial# 0000000002



GTIN 00368071220896  
Serial# 0000000002  
Exp. Date 00-00  
LOT#: 00000

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

## LORATADINE NON DROWSY

loratadine tablet

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-2208(NDC:69230-317)
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<b>Route of Administration</b>	ORAL			
<b>Active Ingredient/Active Moiety</b>				
<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>		
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg		
<b>Inactive Ingredients</b>				
<b>Ingredient Name</b>	<b>Strength</b>			
MAGNESIUM STEARATE (UNII: 70097M6I30)				
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)				
<b>Product Characteristics</b>				
<b>Color</b>	white	<b>Score</b>	no score	
<b>Shape</b>	ROUND	<b>Size</b>	6mm	
<b>Flavor</b>		<b>Imprint Code</b>	439	
<b>Contains</b>				
<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:68071-2208-7	1 in 1 BOX	07/06/2020	
1		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
ANDA	ANDA075209	12/27/2019		

**Labeler** - NuCare Pharmaceuticals,Inc. (010632300)

## Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals,Inc.		010632300	repack(68071-2208)