

LORATADINE ALLERGY RELIEF- loratadine tablet
NuCare Pharmaceuticals, Inc.

Drug Facts

ACTIVE INGREDIENT(S)

Loratadine USP, 10 mg

PURPOSE

Antihistamine

USE(S)

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.

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 and kidney disease	ask a doctor

OTHER INFORMATION

- store between 20 and 25° C (68 and 77° F)
- protect from excessive moisture
- **TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL IS BROKEN OR MISSING FROM BOTTLE.**

INACTIVE INGREDIENTS

Corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

QUESTIONS?

Call **1-800-406-7984**

PRINCIPAL DISPLAY PANEL -

 NuCare Pharmaceuticals, Inc.

NDC: 68071-4788-1

Loratadine 10mg

#100 Tablets

See manufacturer's label for full list of ingredients.

Product #: R0824100

Manufactured by:
Ohm Laboratories, Inc. New Brunswick, NJ 08901

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

Take _____ times a day.
every _____ hours

Rev 01/01/19

WARNING: KEEP OUT OF REACH OF CHILDREN

Loratadine 10mg
Lot: 000000 NDC: 68071-4788-01
MFR NDC: 51660-526-01 Exp.: 00-00

Loratadine 10mg
Lot: 000000 NDC: 68071-4788-01
MFR NDC: 51660-526-01 Exp.: 00-00

GTIN 00368071478815
Serial# 00000000002
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.

STORE AT CONTROLLED TEMPERATURE 59-86°F.

LORATADINE ALLERGY RELIEF

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-4788(NDC:51660-526)
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Route of Administration	ORAL
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Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg

Inactive Ingredients	
Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics			
Color	white (White to Off White)	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	RX526
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-4788-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076134	11/01/2017	

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals,Inc.		010632300	relabel(68071-4788)

Revised: 2/2021

NuCare Pharmaceuticals,Inc.