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 -

Distributed t Ohm Lal Brunswi Packaged B NuCare Pi NuCare Pi Orange, C Orange, C	NDC: 68071-5049-9 Loratadine 10mg	Loratadine 10mg Lot: 000000 NDC: 68071-5049-05 MFR NDC: 51660-526-53 Exp.: 00-00 Serial# 0000000002
a 6 8 0 7 1 5 0 4 9 9 6 Laboratories Inc. New swick, NJ 08901 swick, NJ 08901 dBy: a Pharmaceuticals, Inc. a a Pharmaceuticals, Inc. a a, CA 92867 b instructorsa overy hours every day. bours	#90 Tablets Each tablet contains. Loratadine, USP 10mgAnthistamine 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 drowsness. Stop use and ask a doctor if an allergic reaction to this product	Loratadine 10mg Lot: 000000 NDC: 68071-5049-09 MFR NDC: 51660-526-53 Exp.: 00-00 Serial# 0000000002 GTIN 00368071504996 Serial# 0000000002 Exp. Date 00-00 LOT#: 000000
	occurs. Seek medical help right away. If pregnant or breast-feeding, ask a health professional before use Round White Tablet Debossed: "RX 526" on one side Product #: P0653090	Call your doctor for medical advice about side effects. You may report side effects to FDA at 1-800-FDA-1088.

LORATADINE ALLERGY RELIEF

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-5049(NDC:51660-526)
Route of Administration	ORAL		

Active Ingre	dient/Active Moiety		
	Ingredient Name	Basis of	Strength Strength
	NII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3B		10 mg
			10 mg
Inactive Ing	redients		
	Ingredient Name		Strength
STARCH, CORN	(UNII: 08232NY3SJ)		
LACTOSE MONO	DHYDRATE (UNII: EWQ57Q8I5X)		
MAGNESIUM ST	EARATE (UNII: 70097M6I30)		
Product Cha	racteristics		
Color	white (White to Off White)	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	RX526
Contains			
Packaging			
# Item Code	e Package Description	Marketing Start Date	Marketing End Date
1 NDC:68071- 5049-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	on 09/05/2019	
Marketing	g Information		
Marketing Category		raph Marketing Sta Date	rt Marketing End Date
ANDA	ANDA076134	11/01/2017	

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment					
Name	Address	ID/FEI	Business Operations		
NuCare Pharmaceuticals, Inc.		010632300	repack(68071-5049)		

Revised: 2/2021

NuCare Pharmaceuticals, Inc.