ALLERGY RELIEF- loratadine tablet
NuCare Pharmaceuticals,Inc.

Active ingredient (in each tablet)

Loratadine 10mg

Purpose

Antihistamine

Uses

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

  -runny nose

  -sneezing

  -itching of the nose and throat

  -itchy, watery eyes

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 12 years and over	1 tablet daily; not more than 1 tablet in 24 hours
children under 12 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- **TAMPER EVIDENT:** Do not use if imprinted seal under cap is missing or broken
- store between 2˚C and 30˚C (36˚F-86˚F)
- product of Canada
- protect from excessive moisture

Inactive ingredients

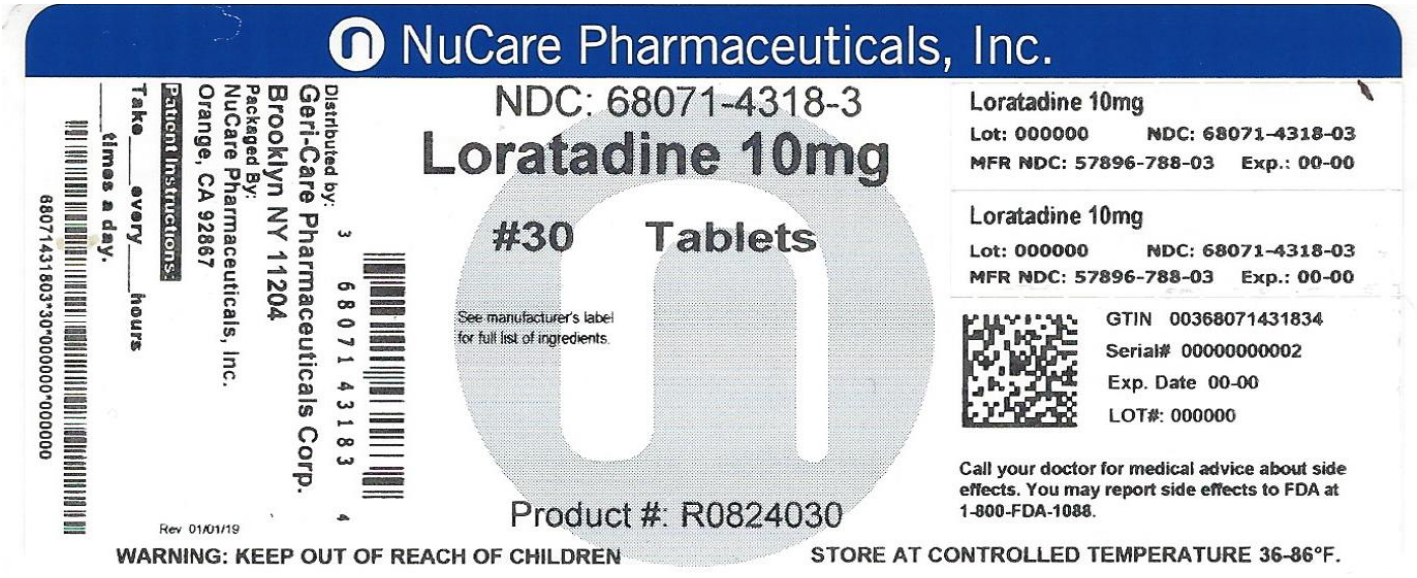
colloidal silicon dioxide, croscarmellose sodium, lactose monohydrate,

magnesium stearate, microcrystalline cellulose

Questions?

call 1-800-540-3765

package label



ALLERGY RELIEF

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68071-4318(NDC:57896-788)
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	

Product Characteristics

Color	white	**Score**	no score
Shape	OVAL	**Size**	8mm
Flavor		**Imprint Code**	LOR;10;APO
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-4318-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	03/07/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076471	01/01/2008	

Labeler - NuCare Pharmaceuticals,Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharamceuticals,Inc.		010632300	relabel(68071-4318)

Revised: 2/2021 NuCare Pharmaceuticals,Inc.