

**LORATADINE ALLERGY RELIEF- loratadine tablet**  
**Lake Erie Medical DBA Quality Care Products LLC**

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**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

Packaged and Distributed by *Quality Care Products, LLC* 1-800-337-8603



Holland, OH 43528

**ALLERGY RELIEF LORATADINE  
10 MG**

Warning: Keep out of children's reach.  
Store between 68 to 77 degrees F.  
Consult with a physician.



WHITE, ROUND, RX;526

GTIN: 00355700097905

NDC: 55700-0097-90

**#90 TABLETS**

Serial:

Lot:

Antihistamine

EXP: //

Each tablet contains Loratadine

Mfr by: Ohm Laboratories Inc., North Brunswick, NJ 08902



## LORATADINE ALLERGY RELIEF

loratadine tablet

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55700-097(NDC:51660-526)
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
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

### Product Characteristics

<b>Color</b>	white (White to Off White)	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	6mm
<b>Flavor</b>		<b>Imprint Code</b>	RX526
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55700-097-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	01/31/2014	
2	NDC:55700-097-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	01/31/2014	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076134	08/28/2003	

**Labeler** - Lake Erie Medical DBA Quality Care Products LLC (831276758)

### Establishment

Name	Address	ID/FEI	Business Operations
Lake Erie Medical DBA Quality Care Products LLC		831276758	repack(55700-097)

Revised: 6/2019

Lake Erie Medical DBA Quality Care Products LLC