

**ALLERGY RELIEF- loratadine tablet, orally disintegrating**  
**Ohm Laboratories 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
- 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 (1-800-222-1222).

## ***DIRECTIONS***

- place 1 tablet on tongue; tablet disintegrates, with or without water

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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

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## ***OTHER INFORMATION***

- Phenylketonurics: Contains Phenylalanine 0.6 mg Per Tablet.
- **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.**
- store between 20° to 25° C (68° to 77° F). Protect from excessive moisture.
- use tablet immediately after opening individual blister.

## ***INACTIVE INGREDIENTS***

Aspartame, croscarmellose sodium, magnesium stearate, mannitol, mint flavor, sodium stearyl fumarate, strawberry cream flavor, tutti-frutti flavor

## ***QUESTIONS?***

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

Follow these directions carefully.  
Do not attempt to push the tablet through the foil.



1. Tear blisters at perforations to separate.



2. Peel printed backing off. Gently remove tablet.



3. Place the tablet on tongue and close mouth. The tablet will dissolve.

## **PRINCIPAL DISPLAY PANEL**

†Compare To the active ingredient of Claritin® RediTabs®

**NDC 51660-527-31**

**ohm®**

**Original Prescription Strength**

***NON-DROWSY\****

**Ages 6 years and older**

**Loratadine Orally Disintegrating Tablets USP, 10 mg**

**ANTIHIISTAMINE**

**Allergy Relief**

**Indoor & Outdoor Allergies**

***No Water Needed***

- ***Melts in Your Mouth***

**24 HOUR**

**Relief of:**

- ***Sneezing***
- ***Itchy, Watery Eyes***
- ***Runny Nose***
- ***Itchy Throat or Nose***

**30 Orally Disintegrating Tablets**

**\*When taken as directed. See Drug Facts Panel.**

**Distributed by: Ohm Laboratories Inc.**

**5217818/R0321**



## ALLERGY RELIEF

loratadine tablet, orally disintegrating

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51660-527
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
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<b>MANNITOL</b> (UNII: 3OWL53L36A)	
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>ASPARTAME</b> (UNII: Z0H242BBR1)	
<b>SODIUM STEARYL FUMARATE</b> (UNII: 7CV7WJK4UI)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	

### Product Characteristics

<b>Color</b>	white (white to off-white)	<b>Score</b>	no score
<b>Shape</b>	ROUND (flat face beveled edged)	<b>Size</b>	10mm
<b>Flavor</b>	STRAWBERRY, TUTTI FRUTTI, MINT	<b>Imprint Code</b>	RC17
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51660-527-31	30 in 1 CARTON	08/31/2007	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077153	08/31/2007	

**Labeler** - Ohm Laboratories Inc. (184769029)

**Registrant** - Ohm Laboratories Inc. (051565745)

### Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	MANUFACTURE(51660-527)

Revised: 4/2021

Ohm Laboratories Inc.