

**LORATADINE- loratadine tablet, orally disintegrating**  
**Walgreen Company**

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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

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**

## **PRINCIPAL DISPLAY PANEL**

**NON-DROWSY\***

**Well at Walgreens**

**NDC 0363-0527-71**

**Wal-itin<sup>®</sup>**

**WALGREENS PHARMACIST RECOMMENDED<sup>‡</sup>**

**24 Hour Allergy Relief**

**Fast Dissolving Tablets**

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

**Antihistamine**

- **Relief of runny nose, sneezing, Itchy throat or nose & Itchy, watery eyes**
- **Ages 6 years & older**

**70 TABLETS**

**INDOOR/OUTDOOR ALLERGIES**

**24 HOUR**

**Compare to Claritin<sup>®</sup> RediTabs<sup>®</sup> active ingredient<sup>‡‡</sup>**

**ORALLY DISINTEGRATING TABLETS**

**\*When taken as directed.**

**See Drug Facts Panel.**

**DISTRIBUTED BY: WALGREEN CO.**

**200 WILMOT RD., DEERFIELD, IL 60015**

**5115100/ORG0315-F**

NON-DROWSY\*

# Wal-itin<sup>®</sup>

24 Hour Allergy Relief  
Fast Dissolving Tablets  
Loratadine Orally Disintegrating  
Tablets USP, 10 mg  
Antihistamine

\* Relief of runny nose, sneezing,  
itchy throat, itchy eyes,  
itchy watery eyes.  
† Ages 6 years & older

**70 TABLETS**  Actual Size

ORALLY DISINTEGRATING TABLETS

**WHEN TAKEN AS DIRECTED, SEE DRUG FACTS PANEL.**



WALGREENS PHARMACY



INDOOR &  
OUTDOOR  
ALLERGIES

Compare to Cl...  
active in

## LORATADINE

loratadine tablet, orally disintegrating

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0527
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
ASPARTAME (UNII: Z0H242BBR1)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UJ)	

**Product Characteristics**

<b>Color</b>	white (White to Off-White)	<b>Score</b>	no score
<b>Shape</b>	ROUND (Flat Faced Beveled Edge)	<b>Size</b>	10mm
<b>Flavor</b>	FRUIT	<b>Imprint Code</b>	RC17
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0527-31	30 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0363-0527-69	10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:0363-0527-71	70 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** - Walgreen Company (008965063)**Registrant** - Ranbaxy Pharmaceuticals Inc. (937890044)**Establishment**

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	MANUFACTURE(0363-0527)

Revised: 3/2015

Walgreen Company