

**CLARITIN REDITABS- loratadine tablet, orally disintegrating**  
**Bayer HealthCare LLC**

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**Claritin**®

**RediTabs**®

**Drug Facts**

**Active ingredient (in each tablet)**

Loratadine 5 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.

**Directions**

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

adults and children 6 years and over	1 tablet every 12 hours; not more than 2 tablets in 24 hours
children under 6 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

### Other information

- safety sealed: do not use if the individual blister unit imprinted with Claritin® RediTabs® is open or torn
- store between 20° to 25°C (68° to 77°F)
- use tablet immediately after opening individual blister
- complies with USP Procedure 2 for Assay and Organic Impurities and Test 2 for Disintegration

### Inactive ingredients

anhydrous citric acid, gelatin, mannitol, mint flavor

### Questions or comments?

**1-800-CLARITIN (1-800-252-7484) or [www.claritin.com](http://www.claritin.com)**

Distributed by MSD Consumer Care, Inc.,  
PO Box 377, Memphis, TN 38151 USA,  
a subsidiary of Merck & Co., Inc., Whitehouse Station, NJ USA.

### PRINCIPAL DISPLAY PANEL - 10 Tablet Carton

NDC 11523-7202-3

***Non-Drowsy\****

***Claritin***®

***RediTabs***®

*loratadine 5 mg/antihistamine*

**Indoor & Outdoor**

**Allergies**

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

**Hour**

**Relief of:**

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

or Nose

**No Water Needed**

**Melts in Your Mouth**

**10**

**ORALLY**

**DISINTEGRATING TABLETS**

**Claritin<sup>®</sup> Reditabs<sup>®</sup>**

**12 Hour**

**Non-Drowsy\***

**Claritin<sup>®</sup>**

**Reditabs<sup>®</sup>**

*loratadine 5 mg/antihistamine*

**Indoor & Outdoor Allergies**

**12 Hour**

No Water Needed  
Melts in Mouth

**BAYER**

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

**ORALLY DISINTE**



**Claritin<sup>®</sup> Reditabs<sup>®</sup>**

**12  
Hour**

**Non-Drowsy\***  
**Claritin<sup>®</sup>**  
**Reditabs<sup>®</sup>**

*loratadine 5 mg/antihistamine*

**Indoor & Outdoor  
Allergies**

**12  
Hour**

**No Water  
Melts in**



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

**ORALLY DISINTE**

## CLARITIN REDITABS

loratadine tablet, orally disintegrating

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:11523-7202
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03B07QN) (LORATADINE - UNII:7AJ03B07QN)	LORATADINE	5 mg

### Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GELATIN (UNII: 2G86QN327L)	
MANNITOL (UNII: 3OWL53L36A)	

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	ROUND (flat faced beveled edge)	<b>Size</b>	12mm
<b>Flavor</b>	MINT	<b>Imprint Code</b>	C5
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-7202-1	3 in 1 CARTON	12/12/2006	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11523-7202-3	1 in 1 CARTON	12/12/2006	
2		10 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
NDA	NDA021993	12/12/2006	

**Labeler** - Bayer HealthCare LLC (112117283)

Revised: 12/2025

Bayer HealthCare LLC