

**LORATADINE ODT - loratadine tablet, orally disintegrating
Amerisource Bergen**

Loratadine Orally Disintegrating Tablets USP 10 mg

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
- sneezing
- itchy, watery eyes
- 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 (1-800-222-1222) right away.

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

Other information

- Phenylketonurics: Contains phenylalanine 2.25 mg per tablet
- do not use if the individual blister unit is open or torn
- store at 20° to 25°C (68° to 77°F)
- use tablet immediately after opening individual blister
- Complies with USP test 2 for Disintegration

Inactive ingredients

aspartame, crospovidone, mannitol, microcrystalline cellulose, peppermint, pregelatinized starch (maize), sodium stearyl fumarate

Questions or comments?

call 1-855-274-4122

Distributed By
AmerisourceBergen
1 West First Avenue
Conshohocken, PA 19428
Questions or Concerns?
www.mygnp.com
Made in India
Code: TS/DRUGS/22/2009

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 10 mg, Blister Carton 10 Orally Disintegrating Tablets

GOOD NEIGHBOR PHARMACY®

Compare to Claritin® RediTabs® active ingredient**

NDC 46122-539-52

Original Prescription Strength

Non-Drowsy*

24 HR

Loratadine Orally Disintegrating

Tablets USP 10 mg

Antihistamine

No Water Needed

Melts in Your Mouth

Indoor & Outdoor Allergies

Relief of:

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

*When taken as directed. See Drug Facts Panel.

actual size

10 Orally Disintegrating Tablets

Drug Facts (continued)

Directions
Place 1 tablet on tongue; tablet disintegrates, with or without water.

adults and children	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

Other information
Phenylethanolamine: Contains phenylethanolamine 2.25 mg per tablet.
do not use if the individual blister unit is open or torn.
store at 20° to 25°C (68° to 77°F).
use tablet immediately after opening individual blister.
Complies with USP test 2 for Disintegration.

Inactive ingredients
aspartame, croscollon, mannitol, microcrystalline cellulose, pectin, polyethylene glycol, polyethylene glycol, sodium stearoyl fumarate.

Questions or comments? call 1-855-274-4122

Drug Facts

Active ingredient (in each tablet)
Loratadine USP 10 mg. Antihistamine

Purpose
Antihistamine

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 (1-800-222-1222) right away.

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SATISFACTION GUARANTEED

**This product is not manufactured or distributed by Bayer HealthCare LLC, distributor of Claritin® RediTabs®.

ABC#: 10186166

Lot: _____
Exp.: _____

GOOD NEIGHBOR PHARMACY

Loratadine Non-Drowsy* **24 HR**

Orally Disintegrating Tablets USP 10 mg

Antihistamine **No Water Needed** **Melts in Your Mouth**

Indoor & Outdoor Allergies

Relief of: • Sneezing • Runny Nose
• Itchy, Watery Eyes • Itchy Throat or Nose

10 Orally Disintegrating Tablets

*When taken as directed, See Drug Facts Panel.

Compare to Claritin® RediTabs® active ingredient**

NDC 46122-539-52

Loratadine
Orally Disintegrating Tablets USP 10 mg

Instructions for Opening Blister Pack

Do not push the tablet from the back

1. Bend and tear blister at perforation

2. Peel off the foil. Gently push tablet out.

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

Unvarnished Zone (dotted line not for printing)

LORATADINE ODT

loratadine tablet, orally disintegrating

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46122-539
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg
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Inactive Ingredients

Ingredient Name	Strength
ASPARTAME (UNII: Z0H242BBR1)	
CROSPROVIDONE (120 .MU.M) (UNII: 68401960MK)	
MANNITOL (UNII: 3OWL53L36A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
PEPPERMINT (UNII: V95R5KMY2B)	
STARCH, CORN (UNII: O8232NY3SJ)	
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)	

Product Characteristics

Color	WHITE (White to Off-white)	Score	no score
Shape	ROUND (Biconvex)	Size	8mm
Flavor	PEPPERMINT	Imprint Code	K;9
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:46122-539-52	1 in 1 CARTON	07/09/2018	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:46122-539-65	3 in 1 CARTON	08/10/2022	
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
ANDA	ANDA208477	07/09/2018	

Labeler - Amerisource Bergen (007914906)

Registrant - Aurohealth LLC (078728447)

Establishment

Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(46122-539) , MANUFACTURE(46122-539)

