

LORATADINE ODT - loratadine tablet, orally disintegrating
Aurohealth LLC

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:

AUROHEALTH LLC

279 Princeton-Hightstown Road
East Windsor, NJ 08520

Made in India

Code: TS/DRUGS/22/2009

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 10 mg, Blister Carton 60 (6 X 10) Orally Disintegrating Tablets

Healthy Living™

**Original Prescription Strength
Panel.**

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

LORATADINE

orally disintegrating tablets USP 10 mg

Antihistamine

Non-Drowsy

Indoor & Outdoor Allergies

**†Compare
to the Active
Ingredient in
Claritin®
RediTabs®**

24 Hour

- Relief of sneezing, runny nose, itchy, watery eyes & itchy throat or nose
- No water needed — melts in your mouth

ACTUAL SIZE
60 (6 X 10)

Orally Disintegrating
Tablets

GLUE - NO COATING

NO COATING

NO COATING

Drug Facts
Active ingredient (in each tablet) Loratadine USP 10 mg
Purpose Antihistamine
Uses temporarily relieve 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 Poison Control Center (1-800-222-2222) right away.

Drug Facts (continued)
Directions Place 1 tablet on tongue; tablet disintegrates, with or without water.
Adults and children 12 years and over 1 tablet daily; not more than 1 tablet in 24 hours.
Children under 6 years of age ask a doctor.
Geriatric patients ask a doctor.
Other information Contains phenylalanine 2.25 mg per tablet. Do not use if in original blister unit is open or torn. Store at 20° to 25° C (68° to 77° F). Store at 15° to 30° C (59° to 86° F) in original blister. Tablets may be broken in half.
Inactive ingredients piperazine, pregabalin, benzocaine (methyl p-aminobenzoate), sodium stearoyl fumarate, croscarmellose sodium, croscarmellose sodium, croscarmellose sodium, croscarmellose sodium, croscarmellose sodium, croscarmellose sodium.
Questions or comments? call 1-855-274-4122

LORATADINE
 orally disintegrating tablets USP 10 mg
Antihistamine

Healthy Living™

Original Prescription Strength *When taken as directed. See Drug Facts Panel.

LORATADINE
 orally disintegrating tablets USP 10 mg
Antihistamine
 Non-Drowsy*
 Indoor & Outdoor Allergies

†Compare to the Active Ingredient in Claritin® RediTabs®

24 Hour
 • Relief of sneezing, runny nose, itchy, watery eyes & itchy throat or nose
 • No water needed — melts in your mouth

ACTUAL SIZE
 60 (6 X 10)
 Orally Disintegrating Tablets

NO Varnish coating area

LM-4887 P1048252

LOT: EXP:

Instructions for Opening Blister Pack

1. Bend and tear blister over perforation.
2. Peel off the foil, gently push tablet out.
3. Place the tablet on tongue and allow to melt in your mouth. Do not swallow the tablet from the pack.

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

Distributed by:
AUROHEALTH LLC
 279 Princeton-Hightstown Road
 East Windsor, NJ 08520
MADE IN INDIA
 Code: TS/DRUGS/22/2009



Healthy Living Loratadine Orally Disintegrating Tablets USP 10 mg, 60 Count

LORATADINE ODT
 loratadine tablet, orally disintegrating

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:58602-881
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)	
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:58602-881-15	6 in 1 CARTON	06/18/2022	
1		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	06/18/2022	

Labeler - Aurohealth LLC (078728447)

Establishment

Name	Address	ID/FEI	Business Operations
Aurobindo Pharma Limited		650381903	ANALYSIS(58602-881) , MANUFACTURE(58602-881)

