# LORATADINE- loratadine tablet, orally disintegrating Dr. Reddy's Laboratories Inc.

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## **Loratidine Orally Disintegrating Tablets, USP**

**Drug Facts** 

## **Active ingredient (in each tablet)**

Loratadine 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.

#### **Directions**

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

adults and children 6 years	1 tablet daily; not more than
and over	1 tablet in 24 hours

children under 6 years of	ask a doctor
age	
consumers with liver or	ask a doctor
kidney disease	

#### Other information

- safety sealed: do not use if the individual blister unit imprinted with Loratadine Orally Disintegrating Tablet, USP 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, mannitol, peppermint flavor, polysorbate 80, pullulan

#### Questions or comments?

call toll-free weekdays 9 AM to 8 PM EST at 1-888-375-3784

Distributed by:

Dr. Reddy's Laboratories Inc.,

Princeton, NJ 08540

Made in India

#### PACKAGE LABEL.PRINCIPAL DISPLAY PANEL

Non-Drowsy\*

## Loratadine Orally Disintegrating Tablets USP, 10 mg

antihistamine

Indoor & Outdoor

## **Allergies**

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

#### 24 Hour Relief of:

Sneezing

Runny Nose

Itchy, Watery Eyes

Itchy Throat or Nose

No Water Needed

#### 30

#### **ORALLY**

#### DISINTEGRATING TABLETS



#### LORATADINE

loratadine tablet, orally disintegrating

#### **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43598-707
Poute of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg	

Inactive Ingredients			
Ingredient Name	Strength		
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)			
PULLULAN (UNII: 8ZQ0AYU1TT)			
POLYSORBATE 80 (UNII: 60ZP39ZG8H)			
MANNITOL (UNII: 30WL53L36A)			

Product Characteristics			
Color	white	Score	no score
Shape	ROUND	Size	12mm
Flavor	MINT	Imprint Code	T10
Contains			

P	Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:43598- 707-30	3 in 1 CARTON	07/01/2023			
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	ANDA213294	05/19/2021	

## **Labeler -** Dr. Reddy's Laboratories Inc. (802315887)

Revised: 6/2023 Dr. Reddy's Laboratories Inc.