# **LORATADINE ODT** - loratadine tablet, orally disintegrating Wal-Mart Stores, Inc.

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## 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?**

1-888-287-1915

PACKAGE LABEL-PRINCIPAL DISPLAY PANEL - 10 mg, Blister Carton 10 (1 x 10) Orally Disintegrating Tablets

NDC 49035-806-03 equate<sup>TM</sup> Compare to Claritin<sup>®</sup> RediTabs<sup>®</sup> active Ingredient\*\* Children's Non-Drowsy\* LORATADINE ORALLY DISINTEGRATING Tablets USP 10 mg Antihis tamine FOR CHILDREN SIX YEARS OF AGE OR OLDER Indoor & Outdoor Allergies LASTS UP TO 24 HOURS Relief of:

SneezingDissolves inRunny NoseDissolves inItchy, Watery EyesYour MouthItchy Throat or Nose\*\*When taken as directed.OriginalSee Drug Facts Panel.Prescription Strength10 (1X10) Orally Disintegrating Tablets



LORATADINE ODT					
loratadine tablet, orally disintegrating					
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Product Information					
Product T ype	HUMAN OTC DRUG	Item Code (Sourc	e)	NDC:4903	5-806
Route of Administration	ORAL				
Active Ingredient/Active Moi	ety				
Ing	gredient Name		Basis of S	trength	Strength
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)			LORATADIN	Ξ	10 mg
Inactive Ingredients					
	Ingredient Name			S	Strength

ASE				
	PARTAME (UNII: ZO			
CRO	<b>DSPOVIDONE (15</b>	<b>MPA.S AT 5%)</b> (UNII: 68401960MK)		
	NNITOL (UNII: 30)	·		
MIC	ROCRYSTALLIN	E CELLULOSE (UNII: OP1R32D61U)		
PEP	PERMINT (UNII: V	95R5KMY2B)		
STA	ARCH, CORN (UNII	: O8232NY3SJ)		
SOI	DIUM STEARYL F	U <b>MARATE</b> (UNII: 7CV7WJK4UI)		
Pre	oduct Characte	ristics		
Col	or WHITE (White to Off-white) Score		no score	
Sha	ape ROUND (Biconvex) Size		8 mm	
Fla	avor PEPPERMINT Imprint Code		К;9	
Car	ntains			
COL	Itallis			
	ckaging			
		Package Description	Marketing Start Date	Marketing End Date
Pa #	ckaging Item Code IDC:49035-806-	Package Description 1 in 1 CARTON	Marketing Start Date	Marketing End Date
<b>Pa</b> #	ckaging Item Code IDC:49035-806-	<b>5</b>		Marketing End Date
<b>Pa</b> # 1 <sup>N</sup> <sub>0</sub>	ckaging Item Code IDC:49035-806-	1 in 1 CARTON 10 in 1 BLISTER PACK; Type 0: Not a Combination		Marketing End Date
<b>Pa</b> # 1 <sup>N</sup> <sub>0</sub>	ckaging Item Code IDC:49035-806-	1 in 1 CARTON 10 in 1 BLISTER PACK; Type 0: Not a Combination		Marketing End Date
<b>P →</b> # 1 N 0 1	ckaging Item Code IDC:49035-806-	1 in 1 CARTON 10 in 1 BLISTER PACK; Type 0: Not a Combination Product		Marketing End Date
Pa # 1 <sup>N</sup> 0 1	<b>ckaging</b> Item Code IDC:49035-806- 3	1 in 1 CARTON 10 in 1 BLISTER PACK; Type 0: Not a Combination Product		Marketing End Date Marketing End Date
Pa # 1 <sup>N</sup> 0 1	ckaging Item Code DC:49035-806- 3 arketing Infe	1 in 1 CARTON 10 in 1 BLISTER PACK; Type 0: Not a Combination Product	04/11/2018	

Labeler - Wal-Mart Stores, Inc. (051957769)

# Registrant - Aurohealth LLC (078728447)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
Vasudha Pharma Chem Ltd.		725431626	API MANUFACTURE(49035-806)	

Establishment					
Name	Address	ID/FEI	<b>Business Operations</b>		
Aurobindo Pharma Limited		650381903	ANALYSIS(49035-806), MANUFACTURE(49035-806)		

Revised: 6/2019

Wal-Mart Stores, Inc.