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

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 equateTM Compare to Claritin[®] RediTabs[®] 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	DSPOVIDONE (15	MPA.S AT 5%) (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 MARATE (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
Pa #	ckaging Item Code IDC:49035-806-	5		Marketing End Date
Pa # 1 ^N ₀	ckaging Item Code IDC:49035-806-	1 in 1 CARTON 10 in 1 BLISTER PACK; Type 0: Not a Combination		Marketing End Date
Pa # 1 ^N ₀	ckaging Item Code IDC:49035-806-	1 in 1 CARTON 10 in 1 BLISTER PACK; Type 0: Not a Combination		Marketing End Date
P → # 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 ^N 0 1	ckaging 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 ^N 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	Business Operations	
Vasudha Pharma Chem Ltd.		725431626	API MANUFACTURE(49035-806)	

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

Revised: 6/2019

Wal-Mart Stores, Inc.