LORATADINE - loratadine tablet
State of Florida DOH Central Pharmacy

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
- itchy, watery eyes
- sneezing
- itching of the nose or throat

How supplied

They are supplied by **State of Florida DOH Central Pharmacy** as follows:

NDC	Strength	Quantity/Form	Color	Source Prod. Code
53808-0457-1	10 mg	30 Tablets in a Blister Pack	WHITE	45802-0650

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

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

- Safety sealed: do not use if the imprinted bottle seal is open or torn.
- Store at 20°-25°C (68°-77°F) (see USP Controlled Room Temperature).

Inactive Ingredients

Lactose monohydrate, magnesium stearate, microcrystalline cellulose and sodium starch glycolate.

Questions or comments?

1 800 719-9260

This product is manufactured by:

Perrigo Company
515 Eastern Avenue Allegan
Michigan 49010

This Product was Repackaged By:

State of Florida DOH Central Pharmacy
104-2 Hamilton Park Drive
Tallahassee, FL 32304
United States

10 mg Label

NDC 53808-0457-1 Non-Drowsy*

LORAtadine

Tablets, USP

10 mg

Antihistamine

Indoor & Outdoor Allergies

24 Hour

Relief of:

• Sneezing

• Runny Nose

• Itchy, Watery

Eyes

• Itchy Throat

or Nose

* When taken as directed.

See Drug Facts Panel.

LORATADINE

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:53808-0457(NDC:45802-0650)
Route of Administration	ORAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics			
Color	WHITE	**Score**	no score
Shape	OVAL	**Size**	6mm
Flavor		**Imprint Code**	L612
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53808-0457-1	30 in 1 BLISTER PACK		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076301	07/01/2009	

Labeler - State of Florida DOH Central Pharmacy (829348114)

Establishment

Name	Address	ID/FEI	Business Operations
State of Florida DOH Central Pharmacy		829348114	repack

Revised: 5/2010 State of Florida DOH Central Pharmacy