LORATADINE- loratadine tablet
HealthMart

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

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. (1-800-222-1222)

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 printed foil under cap is broken or missing. (for bottle only)
- Safety sealed: do not use if the imprinted blister unit is open or torn. (for blister package only)

- Store at 20° to 25°C (68° to 77°F) [see USP Controlled Room Temperature].
- Protect from excessive moisture. (blister unit only)

Inactive Ingredients

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

Questions or comments?

1-800-206-7821

01-2016M

Distributed by McKesson

One Post Street, San Francisco, CA 94104

10 mg Label

HealthMart®

PHARMACY

NDC 62011-0258-2

compare to Claritin®

active ingredient**

loratadine tablets, USP 10 mg

antihistamine

indoor & outdoor allergies

24 hour relief of sneezing/runny nose/

itchy, watery eyes/itchy throat or nose

non-drowsy* • 24 hour

30 tablets

*WHEN TAKEN AS DIRECTED. SEE DRUG FACTS PANEL.

Safety sealed: do not use if printed foil under cap is broken or missing.

LORATADINE

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:62011-0258
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)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics			
Color	WHITE (white to off white)	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	GG296
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:62011-0258-2	1 in 1 CARTON	06/01/2016	05/31/2020
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:62011-0258-3	1 in 1 CARTON	06/01/2016	10/31/2019
2		90 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:62011-0258-4	1 in 1 CARTON	06/01/2016	09/30/2019
3		300 in 1 BOTTLE; Type 0: Not a Combination Product		

4	NDC:62011-0258-1	1 in 1 CARTON	06/01/2016	10/31/2020
4		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	ANDA075209	06/01/2016	10/31/2020

Labeler - HealthMart (177667227)

Revised: 2/2019 HealthMart