LORATADINE- loratadine tablet
CVS 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

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

- store between 20° to 25° C (68° to 77° F)
- protect from excessive moisture
- **TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL IS BROKEN OR MISSING FROM BOTTLE.**

INACTIVE INGREDIENTS

Corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

QUESTIONS?

Call **1-800-406-7984**

PRINCIPAL DISPLAY PANEL - 10 mg Tablet Bottle Carton

CVSHealth™

Compare to the active
ingredient in Claritin®†

Indoor & Outdoor Allergies

Original Prescription Strength

Non-Drowsy*

Allergy Relief

LORATADINE TABLETS, USP 10 mg
Antihistamine

24 Hour Relief of:

1. Sneezing
2. Runny nose
3. Itchy, watery eyes
4. Itchy throat or nose

24
HOUR

Actual Bottle Size on Side Panel
Package Contains One Bottle

45 TABLETS

Actual Size

*When taken as directed.
See Drug Facts Panel.

CVSHealth

Indoor & Outdoor Allergies

Original Prescription Strength
Non-Drowsy*
Allergy Relief
24 HOUR

*When taken as directed. See Drug Facts Panel.

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24 HOUR

Actual Bottle Size on Side Panel
Package Contains One Bottle

45 TABLETS

Actual Size

*When taken as directed. See Drug Facts Panel.

Package Contains One Bottle

Actual Size

Distributed by: CVS Pharmacy, Inc.
One CVS Drive, Woonsocket, RI 02895
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V-14192

CVS Quality Money Back Guarantee

#955066

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R0316

Batch No.

Expiration Date

NON VARNISH

5127715

Drug Facts

Active ingredient (in each tablet)	Purpose
Loratadine, USP 10 mg	Antihistamine

Drug Facts (continued)

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

oor Allergies

LORATADINE

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:59779-528
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**
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	white (White to Off-White)	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	RX526
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-528-56	1 in 1 CARTON	01/08/2010	
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:59779-528-69	1 in 1 CARTON	01/08/2010	
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:59779-528-21	2 in 1 CARTON	01/08/2010	
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:59779-528-31	3 in 1 CARTON	01/08/2010	
4		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:59779-528-43	1 in 1 CARTON	01/08/2010	
5		45 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:59779-528-37	2 in 1 CARTON	01/08/2010	
6		120 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:59779-528-38	1 in 1 CARTON	01/08/2010	
7		365 in 1 BOTTLE; Type 0: Not a Combination Product		
8	NDC:59779-528-60	1 in 1 CARTON	01/08/2010	
8		60 in 1 BOTTLE; Type 0: Not a Combination Product		
9	NDC:59779-528-93	2 in 1 CARTON	01/08/2010	
9		30 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076134	01/08/2010	

Labeler - CVS Pharmacy (062312574)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	MANUFACTURE(59779-528)

Revised: 12/2021

CVS Pharmacy