LORATADINE- loratadine tablet Valu Merchandis ers Company

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.

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 and 25° C (68 and 77° F)
- protect from excessive moisture
- TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.

INACTIVE INGREDIENTS

Corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

QUESTIONS?

Call 1-800-406-7984

PRINCIPAL DISPLAY PANEL - 10 mg Tablet Blister Pack Carton

COMPARE TO THE ACTIVE INGREDIENT IN CLARITIN $^{\circledR}$ *

Best

Choice®

Non-Drowsy[†]

ORIGINAL PRESCRIPTION STRENGTH

ALLERGY RELIEF

Loratadine Tablets, USP 10 mg ANTIHISTAMINE

24 Hour Allergy Relief

Indoor & Outdoor Allergies

Relief of:

- Sneezing
- Itchy, Watery Eyes
- Runny Nose
- Itchy Throat or Nose

10 TABLETS

[†] When taken as directed. See Drug Facts Panel.

Keep the carton. It contains important information. See end panel for expiration date.

Questions? call 1-800-406-7984

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Directions	arom ton which toldet t

get medical help or contact a Poison Control Center right away (1-800-222-1222). Keep out of reach of children. In case of overdose,

Drug Facts (continued)

If pregnant or breast-feeding, ask a health professional

to this product occurs. Seek medical help right away. Stop use and ask a doctor if an allergic reaction

directed. Taking more than directed may cause When using this product do not take more than

disease. Your doctor should determine if you need a Ask a doctor before use if you have liver or kidney

to this product or any of its ingredients. Do not use if you have ever had an allergic reaction Warnings

- itching of the nose or throat

 - ісіну, матегу еуез
 - runny nose
- or other upper respiratory allergies: temporarily relieves these symptoms due to hay fever

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

Drug Facts

Non-Drowsy ORIGINAL PRESCRIPTION STRENGTH

Loratadine Tablets, USP 10 mg

Beșt



R1014

PROUDLY DISTRIBUTED BY: VALU MERCHANDISERS, CO. 5000 KANSAS AVE KANSAS CITY, KS 66106



COMPARE TO THE ACTIVE INGREDIENT IN CLARITIN®

Non-Drowsy[†]

LERGY RELIEF

Loratadine Tablets, USP 10 mg ANTIHISTAMINE

24 Hour Allergy Relief Indoor & Outdoor Allergies Relief of:

Sneezing

Runny Nose

- Itchy, Watery Eyes
- Itchy Throat or Nose

10 TABLETS

† When taken as directed, See Drug Facts Panel,

Non-Drowsy[†] ORIGINAL PRESCRIPTION STRENGTH





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LORATADINE

loratadine tablet

Droduct	Information
PICHICI	111101111111111011

Product Type HUMAN OTC DRUG Item Code (Source) NDC:63941-526

Route of Administration ORAL

Active Ingredient/Active Moiety

2.2021/0 2.108.2021/0 2.202000			
Ingredient Name	Basis of Strength	Strength	
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg	

Inactive Ingredients Ingredient Name Strength STARCH, CORN (UNII: 08232NY3SJ) LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X) MAGNESIUM STEARATE (UNII: 70097M6I30)

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

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63941-526- 69	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	08/19/2003	
2	NDC:63941-526-31	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	08/19/2003	
3	NDC:63941-526- 60	60 in 1 BOTTLE; Type 0: Not a Combination Product	08/19/2003	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076134	08/19/2003	

$\boldsymbol{Labeler} \, \textbf{-} \, \, \text{Valu Merchandisers Company (868703513)}$

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment				
Name	Address	ID/FEI	Business Operations	
Ohm Laboratories Inc.		051565745	manufacture(63941-526)	

Revised: 8/2018 Valu Merchandisers Company