

LORATADINE- loratadine tablet
Chain Drug Marketing Association Inc.

Loratadine

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 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**

Distributed by C.D.M.A., Inc.©
43157 W. Nine Mile
Novi, MI 48376-0995

PRINCIPAL DISPLAY PANEL - 30 Tablet Blister Pack Carton

QUALITY
CHOICE

NDC 63868-151-30

†Compare to
Active Ingredient in
CLARITIN®

Original Prescription Strength

Allergy Relief

Loratadine Tablets, USP
10 mg | Antihistamine

Indoor & Outdoor Allergies

Relief of:
Sneezing | Runny Nose
Itchy, Watery Eyes | Itchy Throat or Nose

24 Hour Allergy Relief | Non-Drowsy*

30 Tablets

*When taken as directed. See Drug Facts Panel.

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

Drug Facts (continued)

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

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5110863



5110863



Allergy Relief

Loratadine Tablets, USP
10 mg | Antihistamine



NDC 63868-151-30

[†]Compare to Active Ingredient in CLARITIN®

Original Prescription Strength

Allergy Relief

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Relief of:
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Allergy Relief

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10 mg | Antihistamine



R0714

Distributed by C.D.M.A., Inc.[®]
43157 W. Nine Mile
Novi, MI 48376-0995
www.qualitychoice.com
Questions: 248-449-9300



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Expiration Date:

Batch No.



Non Varnish Area

LORATADINE

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-151
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	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:63868-151-10	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	08/19/2003	
2	NDC:63868-151-30	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	08/19/2003	
3	NDC:63868-151-01	100 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	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	manufacture(63868-151)

Revised: 8/2018

Chain Drug Marketing Association Inc.