

LORATADINE- loratadine tablet, chewable
KROGER COMPANY

Drug Facts

Active ingredient (in each tablet)

Loratadine USP, 5 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
- itchy, watery eyes
- 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.

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	chew 2 tablets daily; not more than 2 tablets in 24 hours
children 2 to under 6 years of age	chew 1 tablet daily; not more than 1 tablet in 24 hours
children under 2 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- Phenylketonurics: contains phenylalanine 1.25 mg per tablet.
- **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.**

- store between 20° to 25°C (68° and 77°F).

Inactive ingredients

aspartame, citric acid anhydrous, colloidal silicon dioxide, D&C red No. 27 aluminum lake, FD&C blue No. 2 aluminum lake, flavor, magnesium stearate, mannitol, microcrystalline cellulose, sodium starch glycolate, stearic acid

Questions?

call 1-800-406-7984

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

Distributed by:

Ohm Laboratories Inc.

New Brunswick, NJ 08901

0418

Principal Display Panel – 30 Chewable Tablet Blister Pack Carton

NDC 30142-996-30

† Compare to the active ingredient of Children's Claritin® Chewables

ages 2 years and older

Non-Drowsy

Children's Allergy Relief

Loratadine Chewable Tablets USP, 5 mg

Antihistamine

Indoor & Outdoor Allergies

30 CHEWABLE TABLETS

The chewable tablets are to be chewed before swallowing.

Grape Flavored

24 Hour Relief Of:

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

****When taken as directed. See Drug Facts Panel.***



AGES 2 YEARS & OLDER

NON-DROWSY*

Children's Allergy Relief

*When taken as directed. See Drug Facts Panel.

Loratadine Chewable Tablets USP, 5 mg Antihistamine

COMPARE TO the active ingredient of CHILDREN'S CLARITIN® CHEWABLES

**See bottom panel



AGES 2 YEARS & OLDER

NON-DROWSY*

Children's Allergy Relief

NDC 30142-996-30

*When taken as directed. See Drug Facts Panel.

Loratadine Chewable Tablets USP, 5 mg Antihistamine

Indoor & Outdoor Allergies

CHEWABLE

24 Hour Relief of:

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

24 HOUR



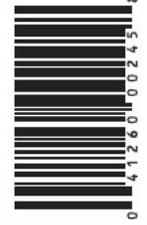
Grape Flavor



actual size

30 CHEWABLE TABLETS

The chewable tablets are to be chewed before swallowing.



0118

DISTRIBUTED BY THE KROGER CO.
CINCINNATI, OHIO 45202
QUALITY GUARANTEE
www.kroger.com

Non Varnish Area

Batch No.

Expiration Date:

**CHILDREN'S CLARITIN® CHEWABLES IS A REGISTERED TRADEMARK OF BAYER HEALTHCARE LLC. BAYER HEALTHCARE LLC IS NOT AFFILIATED WITH KROGER CO. OR THIS PRODUCT.

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Drug Facts (continued)

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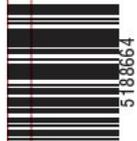
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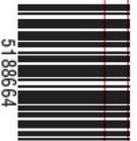
Questions? 1-800-632-6900

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GLUE - NO COATING



5188664



5188664

LORATADINE

loratadine tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30142-996
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	5 mg

Inactive Ingredients

Ingredient Name	Strength
ASPARTAME (UNII: Z0H242BBR1)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
D&C RED NO. 27 (UNII: 2LRS185U6K)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	PURPLE (light purple to dark purple)	Score	no score
Shape	ROUND	Size	10mm
Flavor	GRAPE	Imprint Code	753
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30142-996-30	3 in 1 CARTON	05/22/2019	
1		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	ANDA210088	05/22/2019	

Labeler - KROGER COMPANY (006999528)**Establishment**

Name	Address	ID/FEI	Business Operations
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Ohm Laboratories Inc.

184769029

MANUFACTURE(30142-996)

Revised: 5/2019

KROGER COMPANY