

CHILDRENS LORATADINE- loratadine tablet, chewable
Ohm Laboratories Inc.

Children's Loratadine Chewable Tablets USP, 5 mg

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 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

- chew or crush tablets completely before swallowing.

adults and children 6 years	chew 2 tablets daily; not more than 2 tablets in 24 hours
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and over	
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° to 77°F).

Inactive ingredients

aspartame, carmine, citric acid anhydrous, colloidal silicon dioxide, flavor, magnesium stearate, mannitol, microcrystalline cellulose, sodium starch glycolate, stearic acid

Questions?

call toll-free Monday to Friday 8:30 am to 5:00 pm EST at **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

0619

PRINCIPAL DISPLAY PANEL - 5 mg Tablet Blister Pack Carton

NDC 51660-754-31

**†Compare to the active ingredient of Children's Claritin[®] Chewable
ages 2 years and older**

Children's

Loratadine Chewable Tablets USP, 5 mg

Antihistamine

Indoor & Outdoor Allergies

30 CHEWABLE TABLETS

The chewable tablets are to be chewed before swallowing.

Bubble Gum Flavored

Non-Drowsy*

24 Hour Relief of:

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

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

†Ohm® is a registered trademark of Sun Pharmaceutical Industries, Inc. All other trademarks are property of their respective owners.

ohm[®]
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Drug Facts (continued)

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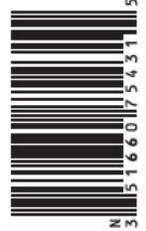
GLUE - NO COATING

Non Varnish Area

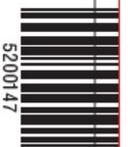
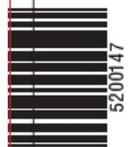
Batch No.

Expiration Date

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R0191
 Distributed by:
 Ohm Laboratories Inc.
 New Brunswick, NJ 08901



loratadine tablet, chewable

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51660-754
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)	
COCHINEAL (UNII: TZ8Z31B35M)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	

Product Characteristics

Color	pink (light to medium pink with a slightly speckled appearance)	Score	no score
Shape	ROUND (Flat face, bevelled edge)	Size	10mm
Flavor	BUBBLE GUM	Imprint Code	754
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51660-754-31	3 in 1 CARTON	06/30/2018	
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	06/30/2018	

Labeler - Ohm Laboratories Inc. (184769029)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(51660-754)

Revised: 1/2024

Ohm Laboratories Inc.