

DYE-FREE LORATADINE- loratadine tablet, chewable
Ohm Laboratories Inc.

Dye-Free Loratadine Chewable Tablets USP, 10 mg

Drug Facts

Active ingredient (in each chewable tablet)

Loratadine USP, 10 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 and over	chew 1 tablets daily; not more than 1 chewable tablets in 24 hours
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children under 6 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, 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

0625

PRINCIPAL DISPLAY PANEL - 10 mg Tablet Blister Pack Carton

Loratadine Chewable Tablets USP, 10 mg
Antihistamine
Indoor & Outdoor Allergies
Dye-Free
24 CHEWABLE TABLETS
The chewable tablets are to be chewed before swallowing.



a SUN PHARMA company

Loratadine Chewable Tablets USP, 10 mg

Antihistamine
Indoor & Outdoor Allergies



24 CHEWABLE TABLETS

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Compare To
the active ingredient of
Claritin® Chewables

NDC 51660-755-02

Dye-Free



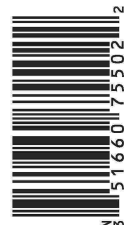
Grape Flavor

Non-Drowsy*

24 Hour Relief of:

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

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



0625

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

Non Varnish Area

Batch No.

Expiration Date:

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Antihistamine
Indoor & Outdoor Allergies

Dye-Free



Grape Flavor

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Drug Facts (continued)

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Inactive ingredients

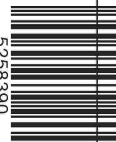
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5258390



DYE-FREE LORATADINE

loratadine tablet, chewable

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51660-755	
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)				
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	white (white to off-white)	Score	no score	
Shape	ROUND (bevelled edge)	Size	10mm	
Flavor	GRAPE	Imprint Code	10	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51660-755-02	3 in 1 CARTON	12/02/2025	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:51660-755-03	6 in 1 CARTON	12/02/2025	
2		8 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	12/02/2025	

Labeler - Ohm Laboratories Inc. (184769029)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		184769029	manufacture(51660-755)

Revised: 1/2026

Ohm Laboratories Inc.