

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

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**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

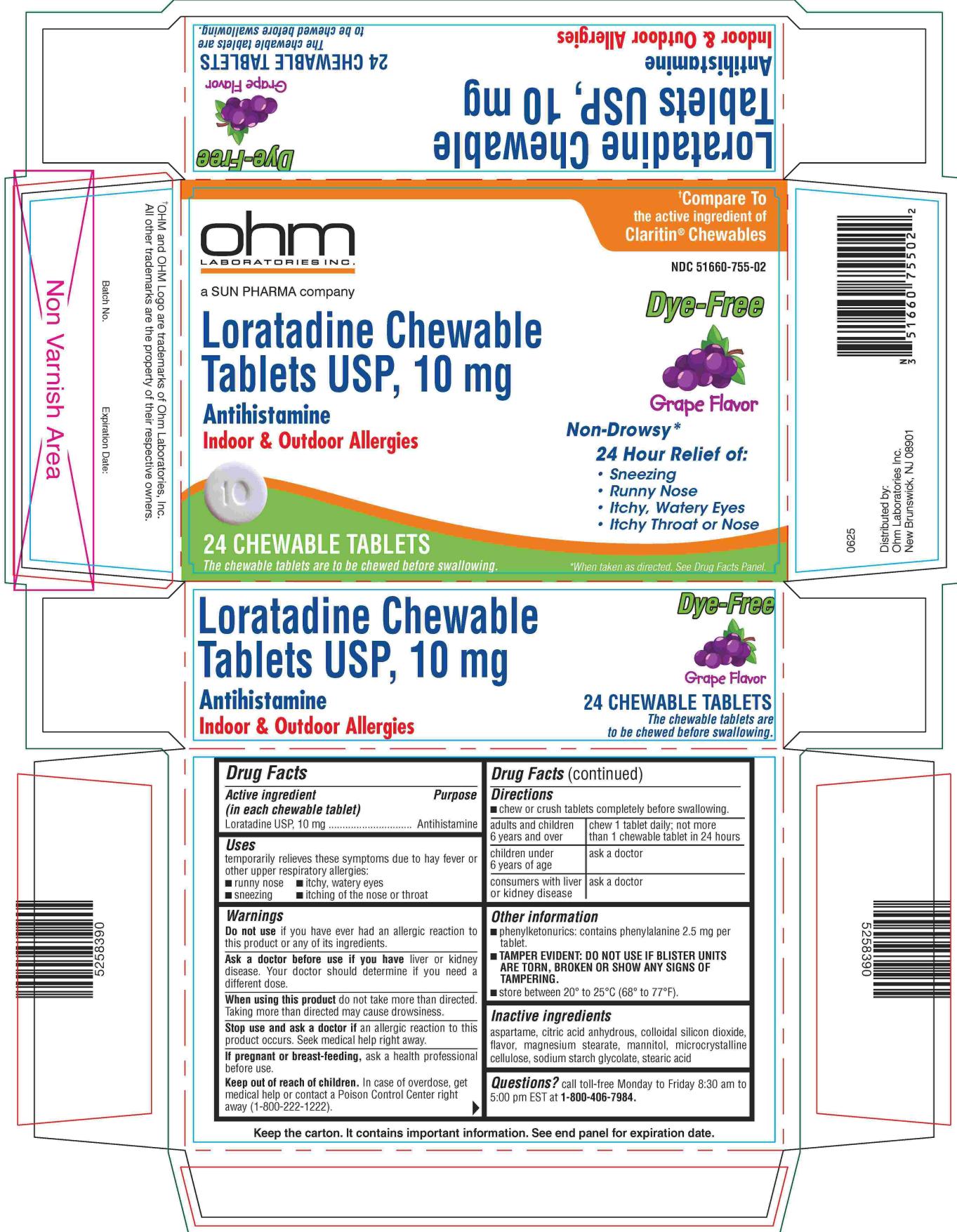
## Questions?

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**



## 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
<b>LORATADINE</b> (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	5 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>ASPARTAME</b> (UNII: Z0H242BBR1)	
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MANNITOL</b> (UNII: 30WL53L36A)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	

## Product Characteristics

<b>Color</b>	white (white to off-white)	<b>Score</b>	no score
<b>Shape</b>	ROUND (bevelled edge)	<b>Size</b>	10mm
<b>Flavor</b>	GRAPE	<b>Imprint Code</b>	10
<b>Contains</b>			

## 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.