LORATADINE- loratadine tablet
Publix Super Markets 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 68° to 77° F (20° to 25° C)
- 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

DISTRIBUTED BY PUBLIX SUPER MARKETS, INC.,
3300 PUBLIX CORPORATE PARKWAY, LAKELAND, FL 33811

PRINCIPAL DISPLAY PANEL - 10 mg Tablet Blister Pack Carton

NDC 56062-526-69

INDOOR & OUTDOOR ALLERGIES
24-HOUR

allergyrelief

LORATADINE TABLETS, USP 10 mg

ANTIHISTAMINE

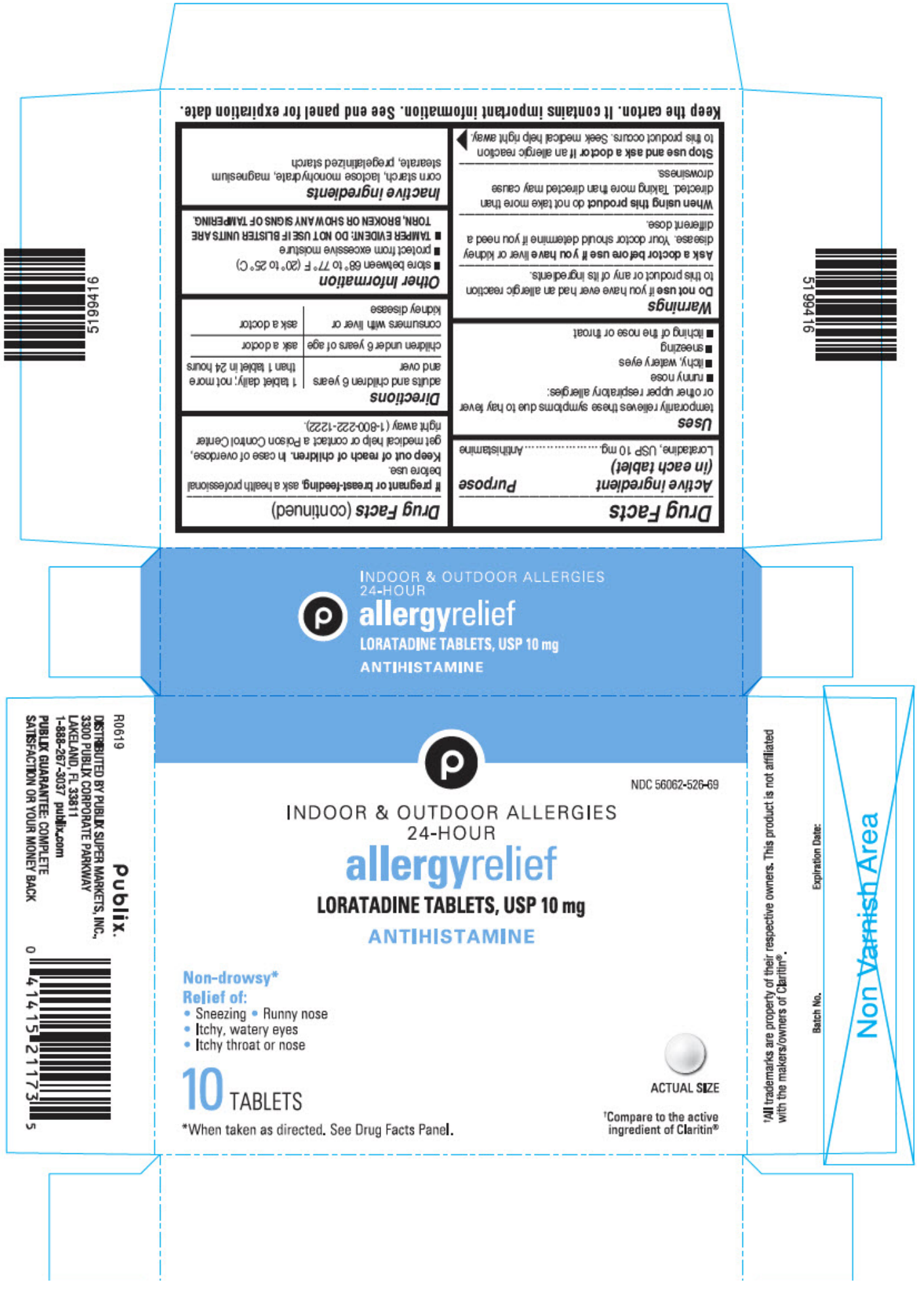
Non-drowsy*
Relief of:
• Sneezing • Runny nose
• Itchy, watery eyes
• Itchy throat or nose

10 TABLETS

*When taken as directed. See Drug Facts Panel.

ACTUAL SIZE

†Compare to the active
ingredient of Claritin®


Keep the carton. It contains important information. See end panel for expiration date.
Drug Facts
Active ingredient (in each tablet)
Purpose
Loratadine, USP 10 mg
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.
Drug Facts (continued)
If pregnant or breast-feeding, ask a health professional before use.
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INDOOR & OUTDOOR ALLERGIES
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ANTIHISTAMINE
Non-drowsy*
Relief of:
• Sneezing • Runny nose
• Itchy, watery eyes
• Itchy throat or nose
10 TABLETS
*When taken as directed. See Drug Facts Panel.
ACTUAL SIZE
†Compare to the active ingredient of Claritin®
†All trademarks are property of their respective owners. This product is not affiliated with the makers/owners of Claritin®.
Batch No.
Expiration Date:
Non Varnish Area
R0619
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LAKELAND, FL 33811
1-888-267-3037 publix.com
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51994 16
51994 16

LORATADINE

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:56062-526
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	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:56062-526-69	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	08/19/2003	
2	NDC:56062-526-31	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	08/19/2003	
3	NDC:56062-526-90	90 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 - Publix Super Markets Inc (006922009)

Registrant - Sun Pharmaceutical Industries Inc. (146974886)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	manufacture(56062-526)

Revised: 9/2019

Publix Super Markets Inc