LORATADINE- loratadine tablet
Cardinal Health 107, LLC

Loratadine Tablets, USP Antihistamine
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:

- itching of the nose or throat
- runny nose
- itchy, watery eyes
- sneezing

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 20° to 25°C (68° to 77°F)
- protect from excessive moisture
- **FOR YOUR PROTECTION:** Do not use if blister is torn or broken.

Overbagged with 10 tablets per bag, NDC 55154-4976-0

WARNING: This Unit Dose package is not child resistant and is Intended for Institutional Use Only. Keep this and all drugs out of the reach of children.

Inactive Ingredients

Corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

Questions?

for Ohm Laboratories call 1-800-406-7984

The drug product contained in this package is from Ohm Laboratories Inc.

Packaged and Distributed by:
American Health Packaging, Columbus, Ohio 43217

Distributed By:

Cardinal Health

Dublin, OH 43017

L4955647-10924 / L4955647-20924

801697

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Principal Display Panel

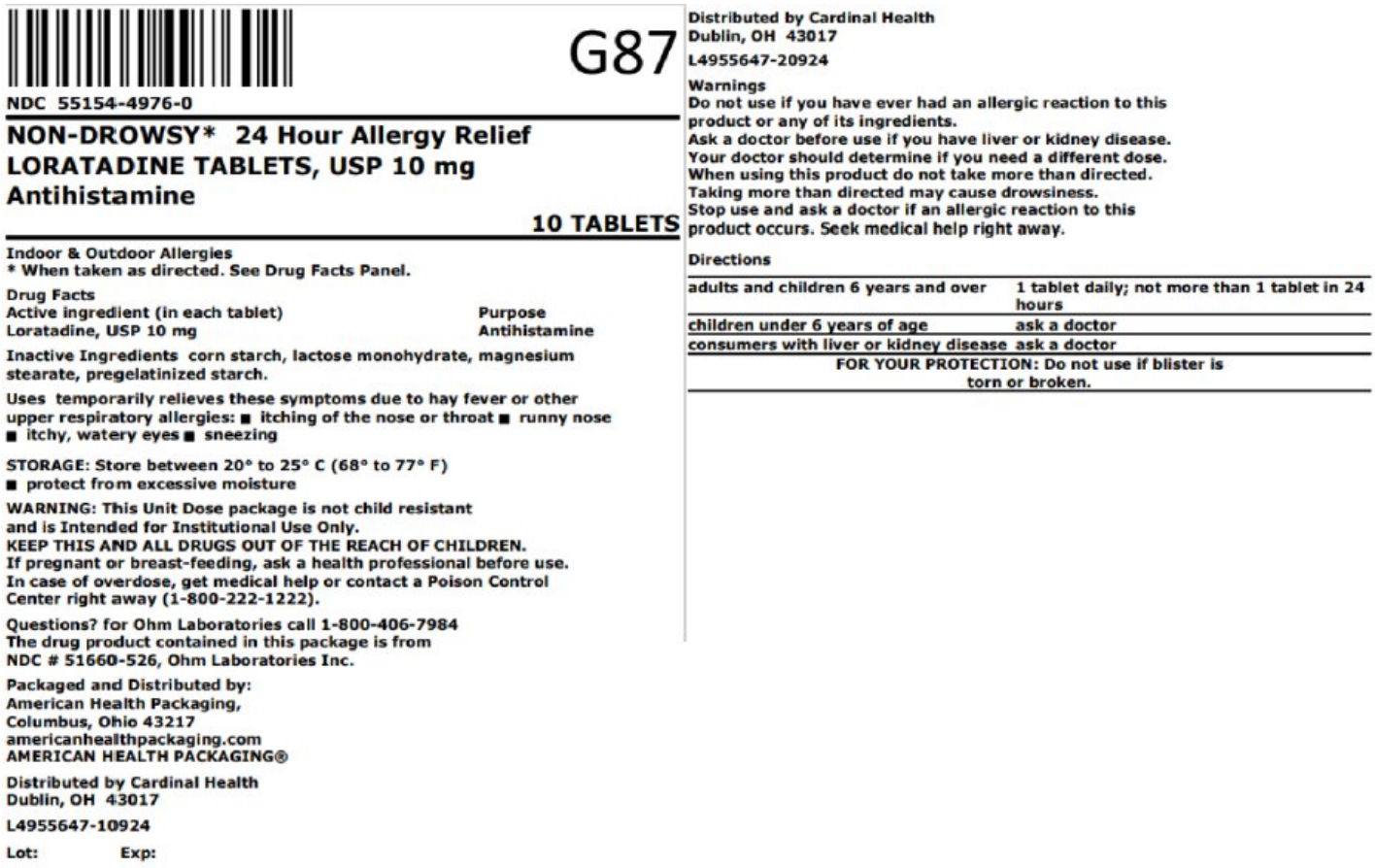
NDC 55154-4976-0

NON-DROWSY* 24 Hour Allergy Relief

LORATADINE TABLETS, USP 10 mg

Antihistamine

10 TABLETS



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NDC 55154-4976-0

NON-DROWSY* 24 Hour Allergy Relief
LORATADINE TABLETS, USP 10 mg
Antihistamine

10 TABLETS

Indoor & Outdoor Allergies
* When taken as directed. See Drug Facts Panel.

Drug Facts

Active ingredient (in each tablet)	Purpose
Loratadine, USP 10 mg	Antihistamine

Inactive Ingredients corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch.

Uses temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: ■ itching of the nose or throat ■ runny nose ■ itchy, watery eyes ■ sneezing

STORAGE: Store between 20° to 25° C (68° to 77° F)
■ protect from excessive moisture

WARNING: This Unit Dose package is not child resistant and is Intended for Institutional Use Only.
KEEP THIS AND ALL DRUGS OUT OF THE REACH OF CHILDREN.
If pregnant or breast-feeding, ask a health professional before use.
In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).

Questions? for Ohm Laboratories call 1-800-406-7984
The drug product contained in this package is from NDC # 51660-526, Ohm Laboratories Inc.

Packaged and Distributed by:
American Health Packaging,
Columbus, Ohio 43217
americanhealthpackaging.com
AMERICAN HEALTH PACKAGING®

Distributed by Cardinal Health
Dublin, OH 43017

L4955647-10924

Lot: Exp:

Distributed by Cardinal Health
Dublin, OH 43017

L4955647-20924

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.

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

FOR YOUR PROTECTION: Do not use if blister is torn or broken.

LORATADINE

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55154-4976(NDC:68084-248)
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:55154-4976-0	10 in 1 BAG	06/14/2013	
1		1 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	ANDA076134	06/14/2013	

Labeler - Cardinal Health 107, LLC (118546603)

Revised: 10/2025 Cardinal Health 107, LLC