LORATADINE- loratadine tablet
Cardinal Health 107, LLC

Drug Facts

*Original Prescription Strength Non-Drowsy**

Indoor and Outdoor Allergies

*When taken as directed. See Drug Facts Panel.

TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.

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.

Directions *(24 Hour Relief)*

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

- **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.**
- **Store at 20° to 25°C (68° to 77°F). [See USP Controlled Room Temperature.]**
- **protect from excessive moisture**

Inactive Ingredients

Corn starch, lactose monohydrate and magnesium stearate.

Questions or comments?

1-800-848-0462

- Serious side effects associated with use of this product may be reported to this number.

Manufactured for:
Mylan Pharmaceuticals Inc.
Morgantown, WV 26505 U.S.A.

Made in India

Code No.: MH/DRUGS/25/NKD/89

Distributed by:
Mylan Institutional Inc.
Rockford, IL 61103 U.S.A.

Distributed by:

Cardinal Health

Dublin, OH 43017

L49557610124

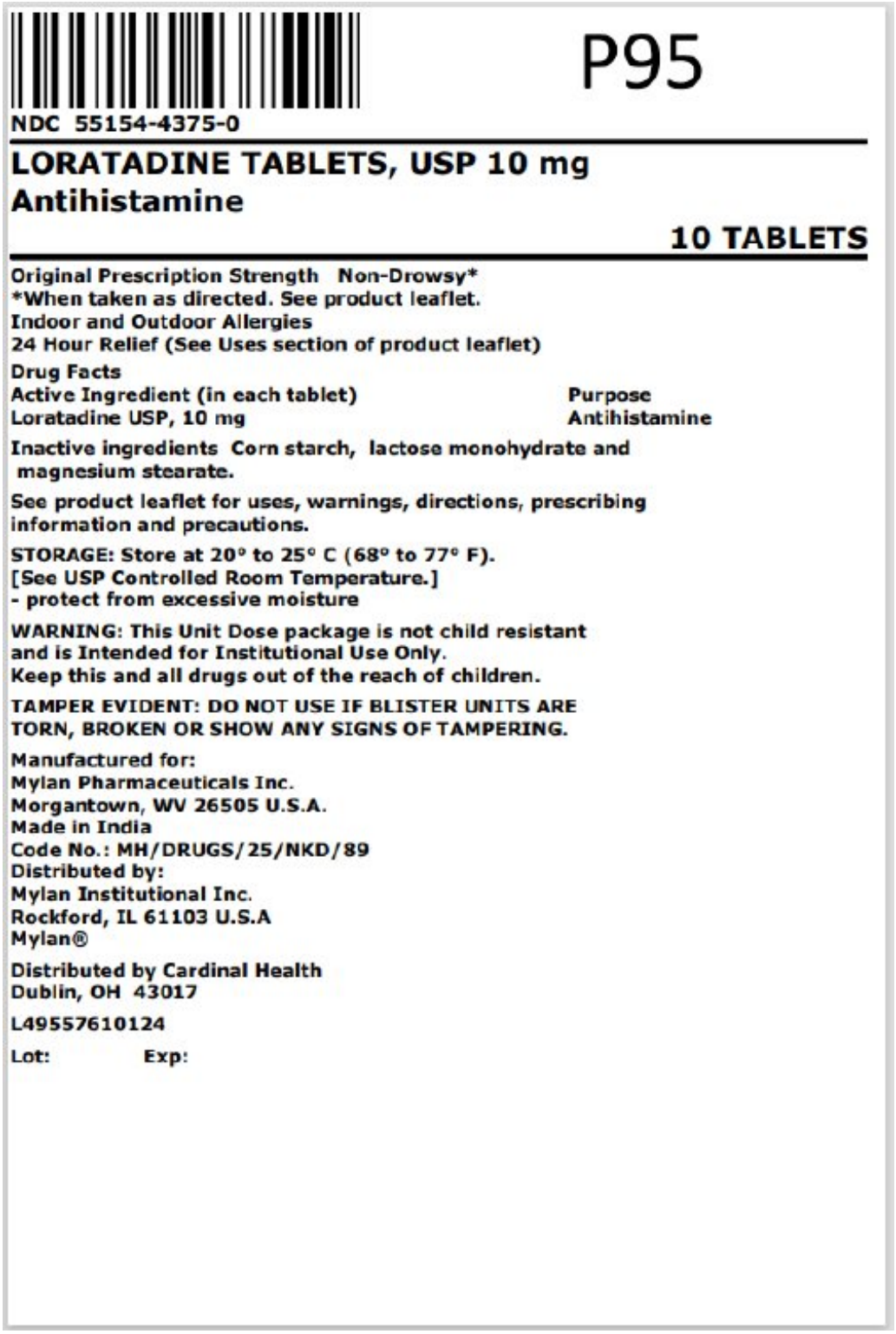
S-11333 R2
11/16

Principal Display Panel

Loratadine Tablets, USP 10 mg

Antihistamine

10 Tablets



P95

NDC 55154-4375-0

LORATADINE TABLETS, USP 10 mg
Antihistamine

10 TABLETS

Original Prescription Strength Non-Drowsy*
***When taken as directed. See product leaflet.**
Indoor and Outdoor Allergies
24 Hour Relief (See Uses section of product leaflet)

Drug Facts

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

Inactive ingredients Corn starch, lactose monohydrate and magnesium stearate.

See product leaflet for uses, warnings, directions, prescribing information and precautions.

STORAGE: Store at 20° to 25° C (68° to 77° F).
[See USP Controlled Room Temperature.]
- 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.

TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.

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L49557610124

Lot: Exp:

LORATADINE

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55154-4375(NDC:51079-246)
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**	G;L;10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55154-4375-0	10 in 1 BAG	04/30/2013	08/31/2024
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	ANDA076154	04/30/2013	08/31/2024

Labeler - Cardinal Health 107, LLC (118546603)

Revised: 1/2024 Cardinal Health 107, LLC