LORATADINE- loratadine tablet, orally disintegrating
Chain Drug Marketing Association Inc.

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

- place 1 tablet on tongue; tablet disintegrates, with or without water

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

- Phenylketonurics: Contains Phenylalanine 0.6 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). Protect from excessive moisture.
- keep in a dry place.
- use tablet immediately after opening individual blister.

INACTIVE INGREDIENTS

Aspartame, croscarmellose sodium, fruit flavors, magnesium stearate, mannitol, sodium stearyl fumarate

QUESTIONS?

Call **1-800-406-7984**

PRINCIPAL DISPLAY PANEL

QC QUALITY CHOICE®

NDC 63868-0157-10

†Compare to the active ingredient in Claritin® Reditabs®

NON-DROWSY■■

Original Prescription Strength

Allergy Relief

Loratadine Orally Disintegrating Tablets, 10 mg

Antihistamine

Indoor & Outdoor Allergies

No Water Needed · Melts in Your Mouth

For Adults and Children six years and older!

Relief of:

Sneezing; Runny Nose

Itchy, Watery Eyes; Itchy Throat or Nose

10 Orally Disintegrating Tablets

24 Hour

Allergy Relief

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

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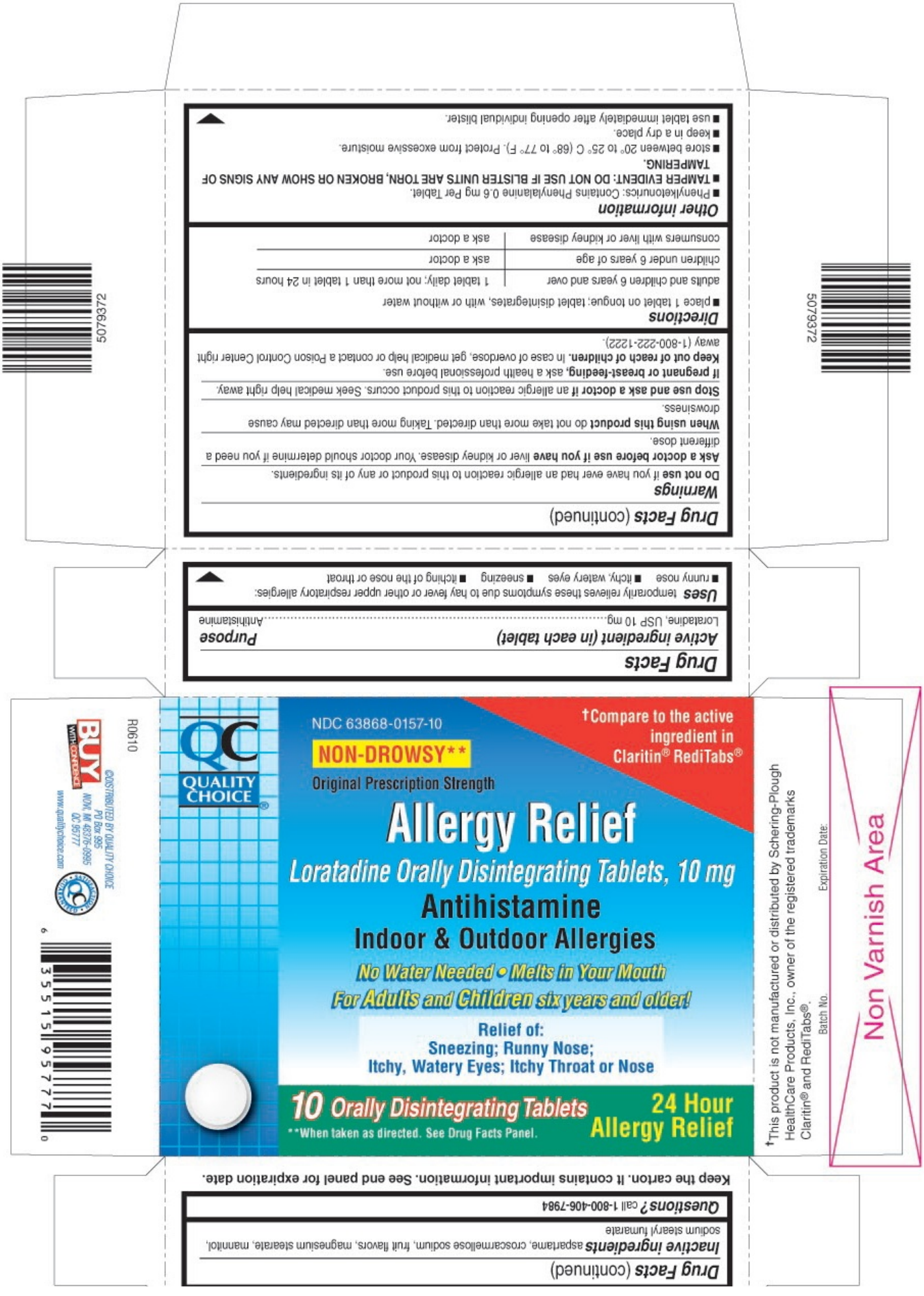
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Itchy, Watery Eyes; Itchy Throat or Nose
10 Orally Disintegrating Tablets
**When taken as directed. See Drug Facts Panel.
24 Hour Allergy Relief
QUALITY CHOICE
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QC 95777
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R0610
†This product is not manufactured or distributed by Schering-Plough HealthCare Products, Inc., owner of the registered trademarks Claritin® and RediTabs®.
Batch No.
Expiration Date:
Non Varnish Area
Drug Facts (continued)
Inactive ingredients aspartame, croscarmellose sodium, fruit flavors, magnesium stearate, mannitol, sodium stearyl fumarate
Questions? call 1-800-406-7984
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
Drug Facts (continued)
Warnings
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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.
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LORATADINE

loratadine tablet, orally disintegrating

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-157
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**
ASPARTAME (UNII: Z0H242BBR1)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MANNITOL (UNII: 3OWL53L36A)	
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)	

Product Characteristics			
Color	white (White to Off-White)	**Score**	no score
Shape	ROUND (flat faced beveled edge)	**Size**	10mm
Flavor	FRUIT	**Imprint Code**	RC17
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63868-157-10	10 in 1 BLISTER PACK		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077153	08/31/2007	

Labeler - Chain Drug Marketing Association Inc. (011920774)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	manufacture(63868-157)