ALLERGY RELIEF- loratadine tablet, orally disintegrating
Cardinal Health

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**

Keep the carton. It contains important information. See end panel for expiration date.

DISTRIBUTED BY

CARDINAL HEALTH

DUBLIN, OHIO 43017

www.myleader.com

1-800-200-6313

PRINCIPAL DISPLAY PANEL

LEADER®

NDC 37205-745-65

Non-Drowsy*

ORIGINAL PRESCRIPTION STRENGTH

Allergy Relief

Loratadine Orally Disintegrating Tablets, 10 mg

Antihistamine

For Adults and Children six years and older!

No Water Needed.

Melts in Your Mouth!

Indoor & Outdoor Allergies

For 24 Hour Relief of:

sneezing; itchy, watery eyes;

runny nose; itchy throat or nose

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

30 ORALLY DISINTEGRATING TABLETS

Compare to Claritin® Reditabs® active ingredient†

This product is not manufactured or distributed by Schering-Plough Healthcare Products, Inc. CLARITIN® and REDITABS® are registered trademarks of Schering Corporation.


LEADER®
Non-Drowsy*
ORIGINAL PRESCRIPTION STRENGTH
Allergy Relief
30 ORALLY DISINTEGRATING TABLETS
Actual Size
SATISFACTION GUARANTEED
NDC 37205-745-65
Compare to Claritin® Reditabs® active ingredient†
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*When taken as directed. See Drug Facts Panel.
†The product is not manufactured or distributed by Schering-Plough Healthcare Products, Inc. CLARITIN® and REDITABS® are registered trademarks of Schering Corporation.
Keep the carton. It contains important information.
See end panel for expiration date.
Batch No.
Expiration Date:
Non Varnish Area
Drug Facts (continued)
■ 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
DISTRIBUTED BY CARDINAL HEALTH
DUBLIN, OHIO 43017
CIN 4596110
www.myleader.com
1-800-200-6313
All Leader® Brand products are 100% satisfaction guaranteed or return to place of purchase for a full refund.
R0612
0 96295 12322 7
5096342
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.
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Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.
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5096342

ALLERGY RELIEF

loratadine tablet, orally disintegrating

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37205-745
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**
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
ASPARTAME (UNII: Z0H242BBR1)	
MANNITOL (UNII: 3OWL53L36A)	
SODIUM STEARYL FUMARATE (UNII: 7CV7WJK4UI)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics			
Color	white (white to off-white)	**Score**	no score
Shape	ROUND (flat faced beveled edge)	**Size**	10mm
Flavor	STRAWBERRY, TUTTI FRUTTI, MINT	**Imprint Code**	RC17
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:37205-745-65	30 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 - Cardinal Health (097537435)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	manufacture(37205-745)