LORATADINE- loratadine tablet Amerisource Bergen

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.

DIRECTIONS

| | 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 and 25° C (68 and 77° F)
- 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

QUESTIONS?

Call 1-800-406-7984

PRINCIPAL DISPLAY PANEL

GOOD NEIGHBOR PHARMACY®

Compare to Claritin® Tablets active ingredient†

NDC 46122-0158-65

Original Prescription Strength

Loratadine Tablets USP, 10 mg

24 HOUR

Non-Drowsy*

Antihis tamine

Indoor & Outdoor Allergies

Relief of:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Throat or Nose

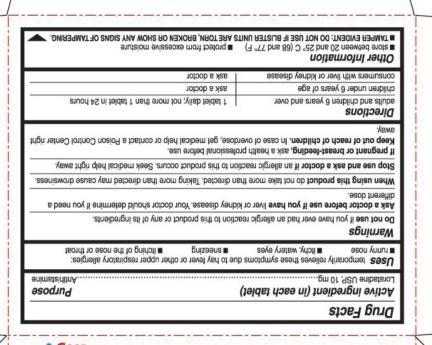
30 tablets

*When taken as directed.

See Drug Facts Panel.

Distributed By AmerisourceBergen

5096598/0612





Loratadine Tablets USP, 10 mg

Antihistamine

Indoor & Outdoor Allergies



Compare to Claritin® Tablets active ingredient

NDC 46122-0158-65

Original Prescription Strength

24 HOUR Non-Drowsy*

Loratadine

Tablets USP, 10 mg

Antihistamine

Indoor & Outdoor Allergies

Relief of:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Throat or Nose

30 tablets

*When taken as directed.

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

Questions? call 1-800-406-7984

Inactive ingredients corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

Drug Facts (continued)

This product is not manufactured or distributed by Schering-Plough HealthCare Products, Inc. CLARITIN® is a registered trademark of Schering Corporation.

Distributed By
AmerisourceBergen
1300 Morris Drive
Chesterbrook, PA 19087
Visit us at
www.goodneighborpharmacy.com

LORATADINE

loratadine tablet

| Produc | t Infor | mation |
|--------|-------------------|--------|
| Promin | 1 I I I I I O F I | |

Product Type HUMAN OTC DRUG Item Code (Source) NDC:46 122-158

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) | | | |
| STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ) | | | |

| Product Characteristics | | | |
|-------------------------|----------------------------|--------------|----------|
| Color | white (White to Off-White) | Score | no score |
| Shape | ROUND | Size | 6 mm |
| Flavor | | Imprint Code | RX526 |
| Contains | | | |

| Packaging | | | |
|--------------------|----------------------|----------------------|--------------------|
| # Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 NDC:46122-158-65 | 30 in 1 BLISTER PACK | | |

| Marketing Information | | | |
|-----------------------|--|----------------------|--------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| ANDA | ANDA076134 | 08/19/2003 | |
| | | | |

Labeler - Amerisource Bergen (007914906)

$\pmb{Registrant - \text{Ranbaxy Pharmaceuticals Inc. (937890044)}}$

Establishment

| Name | Address | ID/FEI | Business Operations |
|-----------------------|---------|-----------|------------------------|
| Ohm Laboratories Inc. | | 051565745 | manufacture(46122-158) |

Revised: 10/2012 Amerisource Bergen