

ALL DAY ALLERGY RELIEF- loratadine tablet
P & L Development, LLC

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
- sneezing
- itchy, watery eyes
- 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

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 at 20°-25°C (68°-77°F) (see UPS Controlled Room Temperature

Inactive ingredients

lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal Display Panel

Compare to the active ingredient in Claritin®†

all day allergy relief

loratadine tablets 10 mg

non-drowsy*

Indoor & Outdoor Allergies

24 hour relief of:

- sneezing
- runny nose
- itchy, watery eyes
- itchy throat or nose

gluten-free

*when taken as directed, see drug facts panel.

tablets

†This product is not manufactured or distributed by Bayer Healthcare LLC, distributor of Claritin®.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOW ANY SIGN OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT

INFORMATION

Distributed by:

PL Developments

200 Hicks Street

Westbury, NY 11590

Package Label

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

Drug Facts (continued)	
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	children under 6 years of age ask a doctor
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Inactive ingredients	lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate
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all day allergy relief

loratadine tablets 10 mg
antihistamine



Compare to the active ingredient in **Claritin**†
NDC 59726-758-10

all day allergy relief

loratadine tablets 10 mg
antihistamine

non-drowsy*
indoor & outdoor allergies

24 hour relief of:

- sneezing
- itchy, watery eyes
- runny nose
- itchy throat or nose

aluten-free

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 KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

green box

*when taken as directed. see drug facts panel.

10 tablets



actual size

TAMPER EVIDENT

Lot No.:
Exp. Date:



Distributed by:
PL Developments
200 Hicks Street
Westbury, NY 11590

PLD-A532A FC005521



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READYinCASE All day allergy relief

ALL DAY ALLERGY RELIEF

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59726-758
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII: 7AJ03BO7QN)	LORATADINE	10 mg

Inactive Ingredients

Ingredient Name	Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	GG296
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59726-758-10	10 in 1 CARTON	09/30/2018	07/26/2024

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	ANDA075209	09/30/2018	07/26/2024

Labeler - P & L Development, LLC (800014821)

Revised: 10/2021

P & L Development, LLC