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

NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING. KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

(1-800-SSS-1SSS): help or contact a Poison Control Center right away keep out of reach of children. In case of overdose, get medical If pregnant or breast-feeding, ask a health professional before occurs. Seek medical help right away. Stop use and ask a doctor if an allergic reaction to this product I aking more than directed may cause drowsiness. when using this product, do not take more than directed. Your doctor should determine if you need a different dose. Ask a doctor before use if you have liver or kidney disease. broduct or any of its ingredients. Do not use if you have ever had an allergic reaction to this sbuuuem пісну, матепу еуев itching of the nose or throat

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or other upper respiratory allergies:

NSGS Temporarily relieves these symptoms due to hay fever

Loratadine, USP 10 mg. Antihistamine

> (10 each tablet) Active ingredient

Purpose

Drug Facts

all day allergy relief

loratadine tablets 10 mg antihistamine



Compare to the active ingredient in Claritin®1 NDC 59726-758-10

all day allergy relief

loratadine tablets 10 mg antihistamine



non-drowsy* indoor & outdoor allergies

24 hour relief of:

 itchy, watery eyes sneezing

Bayer HealthCare LLC, distributor of Claritin®.

Cuestions or comments?

(emperature)

or kidney disease

children under 6

6 years and over

Directions

adults and children

Drug Facts (continued)

years of age

cousnimets with liver

Other information

This product is not manufactured or distributed by

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

atearate, microcrystalline cellulose, sodium starch glycolate

■ store at 20°-25°C (68°-77°F) (see USP Controlled Room

92K & doctor

SSK S GOCTOF

than 1 tablet in 24 hours.

Tablet daily; not more

Inactive ingredients lactose monohydrate, magnesium

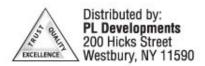
itchy throat or nose

aluten-free

runny nose



Lot No.:





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: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	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