ALLERGY RELIEF- loratadine tablet
QUALITY CHOICE (Chain Drug Marketing Association)

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 (1-800-222-1222) right away.

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 20º-25ºC (68º-77ºF) (see UPS Controlled Room Temperature

Inactive ingredients

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

Questions or comments?

Call **1-248-449-9300** Monday-Friday 9AM-5PM EST

Principal Display Panel

†Compare to the Active Ingredient in Claritin®

Original Prescription Strength

Allergy Relief

Loratadine Tablets, USP

10 mg | Antihistamine

Indoor & Outdoor Allergies

Relief of:

Sneezing | Runny Nose

Itchy, Watery Eyes | Itchy Throat or Nose

Gluten Free

24 Hour Allergy Relief | Non-Drowsy*

Tablets

*When taken as directed. See Drug Facts panel.

†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 C.D.M.A., Inc.©

43157 W 9 Mile Rd

Novi, MI 48375

www.qualitychoice.com

Package Label

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
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- itching of the nose or throat

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Drug Facts (continued)

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 USP Controlled Room Temperature)

Inactive ingredients

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

Questions or comments?

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QUALITY CHOICE

Allergy Relief

Loratadine Tablets, USP, 10 mg | Antihistamine

10 Tablets

NDC 63868-462-10

QUALITY CHOICE

†Compare to the Active Ingredient in Claritin®

Original Prescription Strength

Allergy Relief

Loratadine Tablets, USP 10 mg | Antihistamine

Indoor & Outdoor Allergies

Relief of:

Sneezing | Runny Nose

Itchy, Watery Eyes | Itchy Throat or Nose

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SEE OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

QUALITY CHOICE Allergy Relief

ALLERGY RELIEF

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-462
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:63868-462-10	10 in 1 CARTON	02/28/2019	02/28/2025
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:63868-462-30	1 in 1 BOX	02/28/2019	02/28/2025
2		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:63868-462-01	1 in 1 BOX	02/28/2019	02/28/2025
3		100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075209	02/28/2019	02/28/2025

Labeler - QUALITY CHOICE (Chain Drug Marketing Association) (011920774)

Revised: 11/2022 QUALITY CHOICE (Chain Drug Marketing Association)