

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 20°-25°C (68°-77°F) (see UPS Controlled Room Temperature)
- protect from light

Inactive ingredients

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

Questions or comments?

Call **800-935-2362** Monday-Friday 9AM-5PM EST

Principal Display Panel

†Compare to the Active Ingredient in Claritin® 24 Hour

Original Prescription Strength

Allergy Relief

Loratadine 10 mg Tablets

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® 24 Hour

TAMPER EVIDENT: DO NOT USE IF PRINTED SFETY SEAL UNDER CAP IS BROKEN OR MISSING

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

<p>Drug Facts</p> <p>Active ingredient (in each tablet) Loratadine, USP 10 mg.....Antihistamine</p> <p>Purpose Antihistamine</p> <p>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</p> <p>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.</p> <p><small>This product is not manufactured or distributed by Bayer HealthCare LLC, distributor of Claritin® 24 Hour.</small></p> <p>SATISFACTION GUARANTEED 100% Distributed by C.D.M.A., Inc. © 43157 W 9 Mile Rd Novi, MI 48375 www.qualitychoice.com Questions: 800-935-2362</p>	<p>Drug Facts (continued)</p> <p>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</p> <p>Other information ■ store at 20°-25°C (68°-77°F) (see USP Controlled Room Temperature) ■ protect from light</p> <p>Inactive ingredients lactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate</p> <p>Questions or comments? Call 1-800-935-2362 Monday-Friday 9AM-5PM EST</p> <p>TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.</p> <p>KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.</p>	<p>QC QUALITY CHOICE NDC 63868-414-01</p> <p>*Compare to the Active Ingredient in Claritin® 24 Hour</p> <p>Original Prescription Strength</p> <p>Allergy Relief</p> <p>Loratadine Tablets, 10 mg Antihistamine</p> <p>Indoor & Outdoor Allergies</p> <p>Relief of: Sneezing Runny Nose Itchy, Watery Eyes Itchy Throat or Nose Gluten Free 24 Hour Allergy Relief Non-Drowsy*</p> <p>100 Tablets *When taken as directed. See Drug Facts panel. actual size</p>	<p>QC QUALITY CHOICE NDC 63868-414-01</p> <p>*Compare to the Active Ingredient in Claritin® 24 Hour</p> <p>Original Prescription Strength</p> <p>Allergy Relief</p> <p>Loratadine Tablets, 10 mg Antihistamine</p> <p>Indoor & Outdoor Allergies</p> <p>Relief of: Sneezing Runny Nose Itchy, Watery Eyes Itchy Throat or Nose Gluten Free 24 Hour Allergy Relief Non-Drowsy*</p> <p>100 Tablets *When taken as directed. See Drug Facts panel. actual size</p>
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Item # 99736



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Lot No.:
Exp. Date:

1 BOTTLE INSIDE

QUALITY CHOICE Allergy Relief

ALLERGY RELIEF			
loratadine tablet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-414
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	439	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-414-01	1 in 1 BOX	04/15/2021	04/15/2026
1		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	04/15/2021	04/15/2026	

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

Revised: 12/2025

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