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

†This product is not manufactured or distributed by Bayer Healthcare LLC, distributor of Claritin® 24 Hour

^{*}When taken as directed. See Drug Facts panel.

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



ALLERGY RELIEF

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-414	
Route of Administration	ORAL			
Active Ingredient/Active Moietv				

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	439
Contains			

Packaging				
# Item Code	Package Description	Marketing Start Date	Marketing End Date	
NDC:63868- 414-01	1 in 1 BOX	04/15/2021		
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		

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

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