

CAREALL LORATADINE- loratadine tablet
New World Imports

Drug Facts

Loratadine 10mg

Antihistamine

In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222)

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

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.

Adult and children over 6 years old: 1 tablets daily: no 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

lactose monohydrate, magnesium stearate, sodium starch glycolate. May contain: microcrystalline cellulose, pregelatinized starch

Store between 20-25 degrees celcius (68-77 degrees farhenheit).

Protect from excessive moisture

Do not use if imprinted seal under bottle cap is broken or missing

You may report side effects 1-888-952-0050

NDC 51824-075-01

CAREALL®

Non-Drowsy • 24 Hour Allergy Relief

Antihistamine

RELIEVES

- ▶ Sneezing
- ▶ Itchy, watery eyes
- ▶ Runny nose & itchy throat

30 Loratadine Tablets
10mg each

Compares to the active ingredient in CLARITIN®.

Your Source For High Quality

Drug Facts

Active Ingredient (in each tablet)
Loratadine 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

Drug Facts (continued on inside)
Distributed By: NWI, Inc., 160 Athens Way, Nashville, TN 37228

Made in India
LB1761 R0220

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Lot: Exp:
PEEL HERE

Drug Facts (continued)

medical help right away.

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of accidental overdose, contact a doctor or Poison Control Center (1-800-222-1222) right away.

Directions

Adults and children 6 years and over	1 tablet daily; no 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 room temperature 20° – 25°C (68° – 77°F)
- protect from excessive moisture
- do not use if imprinted seal under safety cap is broken or missing
- You may report side effects 1-888-952-0050

Inactive Ingredients
lactose monohydrate, magnesium stearate, sodium starch glycolate

May contain: microcrystalline cellulose, pregelatinized starch

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

CAREALL LORATADINE

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51824-075
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
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	white (White to off white)	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	GG296;G;10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51824-075-01	30 in 1 BOTTLE; Type 0: Not a Combination Product	03/02/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210722	03/02/2020	

Labeler - New World Imports (075372276)

Registrant - New World Imports (075372276)

Revised: 12/2023

New World Imports