CLARITIN- loratadine tablet
R J General Corporation

Drug Facts

CLARITIN ®

Active ingredients (in each tablet)	***Purpose***
Loratadine 10 mg	Antihistamine

USE

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 productdon not take more than directed. Taking more than directed may cause drowsiness.

Stop use and ask doctor ifan allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding,ask 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.

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

consumer with liver or kidney disease	ask a doctor

Other information

- Safety sealed: do not use if the individual blister unit imprinted with Claritin ® is open or torn
- Store between 20 º to 25 º C(68 ºto 77 ºF)
- Protect from excessive moisture

Inactive ingredients

Corn starch, lactose monohydrate, magnesium stearate

Questions and Comments?

1-800-CLARITIN(1-800-252-7484)or **www.claritin.com**

Dist. by: Bayer HealthCare LLC

Whippany, NJ 07981

Product of India

Do not use if pouch is torn or broken.

Repackaged and distributed by:

RJ General

2024 Northwest Drive

Cincinnati OH 45231

PRINCIPAL DISPLAY PANEL - 25 Tablet Pouch Carton

Non-Drowsy*

Claritin ®

loratadine tablets 10 mg/ antihistamine

Indoor & Outdoor Allergies

Relief of:

- **runny nose**
- **itchy, watery eyes**
- **sneezing**
- **itching of the throat or nose**

CLARITIN

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70264-030(NDC:11523-6655)
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)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	458;Claritin;10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70264-030-01	25 in 1 CARTON	12/26/2022	
1		1 in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA019658	03/01/2021	

Labeler - R J General Corporation (122542830)

Establishment

Name	Address	ID/FEI	Business Operations
R J General Corporation		122542830	repack(70264-030)

Revised: 11/2025 R J General Corporation