CLARITIN- loratadine tablet
Select Consumer Group

Drug Facts

Active ingredient (in each tablet)

Loratadine, 10 mg USP

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

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

When using this product do not take more than directed. Taking more than directed may cause drowsiness.

Stop use

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

If pregnant or breast-feeding

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

Keep out of reach of children

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

Directions

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 -25º C (68º -77º F) (see USP Controlled Room Temperature)

Inactive ingredientslactose monohydrate, magnesium stearate, microcrystalline cellulose, sodium starch glycolate

Questions or comments?

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

Carton label 10 count tablets

Original Prescription Strength

Non-Drowsy*

Claritin®

loratadine tablets 10 mg/antihistamine

Indoor & Outdoor

Allergies

24

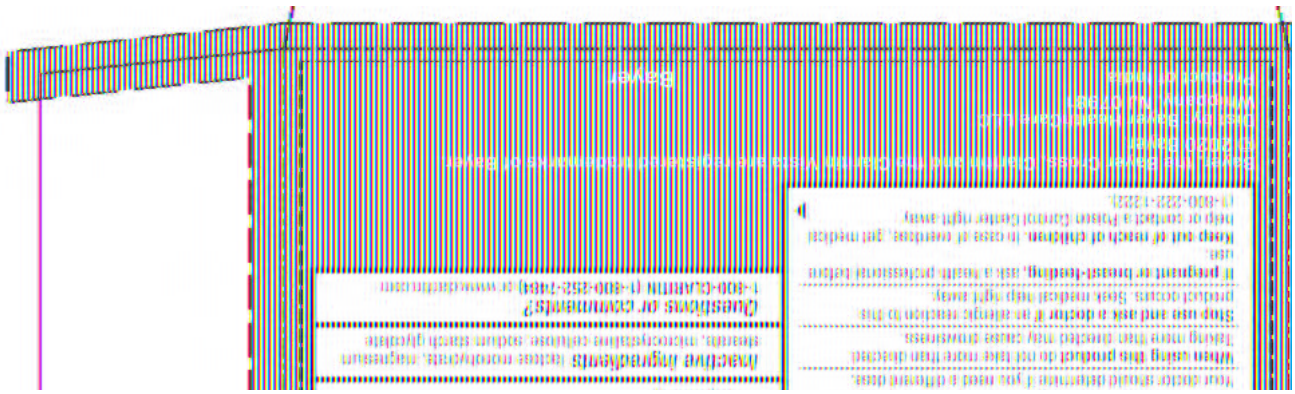
Hour

Relief of:

- ***Sneezing***
- ***Runny Nose***
- ***Itchy, Watery Eyes***
- ***Itchy Throat or Nose***

**When taken as directed.*

See Drug Facts Panel



CLARITIN

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:85237-1630(NDC:11523-0800)

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)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM STARCH GLYCOLATE TYPE A CORN (UNII: AG9B65PV6B)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:85237-1630-1	1 in 1 CARTON	08/18/2025	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075209	12/01/2020	

Labeler - Select Consumer Group (119185084)

Establishment

Name	Address	ID/FEI	Business Operations
Select Consumer Group		119185084	repack(85237-1630)

Revised: 8/2025 Select Consumer Group