LORATADINE- loratadine tablet
Major Pharmaceuticals
Reference Label Set Id: 71271c28-c0dd-4992-a8a9-6d0d57f7498f

Major Pharmaceuticals Loratadine Drug Facts

Active ingredient (in each tablet)

Loratadine 10 mg

Purpose

Antihistamine

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 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 right away. (1-

800-222-1222)

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

- do not use if printed foil under cap is broken or missing
- store between 20˚ to 25˚C (68˚ to 77˚F)

Inactive ingredients

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

Questions or comments?

1-800-719-9260

Principal Display Panel

MAJOR®

compare to the active ingredient in Claritin® Tablets

non-drowsy*

loratadine

loratadine tablets, 10 mg

antihistamine

indoor & outdoor allergies

original prescription strength

24 hour relief of:

• sneezing

• runny nose

• itchy, watery eyes

• itchy throat or nose

actual size

90 tablets

*when taken as directed. see drug facts panel.

LORATADINE

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0904-6852
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)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	

Product Characteristics

Color	WHITE	**Score**	no score
Shape	OVAL	**Size**	8mm
Flavor		**Imprint Code**	L612
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0904-6852-60	1 in 1 CARTON	12/06/2018	11/30/2024
1		100 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0904-6852-89	1 in 1 CARTON	04/08/2019	
2		90 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:0904-6852-07	30 in 1 CARTON	08/01/2019	12/31/2024
3		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:0904-6852-72	1 in 1 CARTON	06/04/2019	
4		300 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:0904-6852-61	100 in 1 CARTON	02/21/2005	01/31/2026
5		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	ANDA076301	02/21/2005	

Labeler - Major Pharmaceuticals (191427277)

Revised: 12/2025 Major Pharmaceuticals