

**LORATADINE- loratadine tablet**  
**Apotex Corp.**

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**Loratadine Tablet 10 mg**

***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
- 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 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***

- safety sealed: do not use if induction seal, with "Lift N Peel" tab, under cap is broken or missing
- store between 2°C and 30°C (36°F and 86°F)
- protect from excessive moisture

### ***Inactive ingredients***

colloidal silicon dioxide, croscarmellose sodium, lactose monohydrate, magnesium stearate, microcrystalline cellulose

### ***Questions or comments?***

Call **1-800-706-5575**, weekdays, 8:30 am - 5:00 pm Eastern Standard Time

Manufactured by: Apotex Inc. Toronto, Ontario Canada M9L 1T9	Manufactured for: Apotex Corp. Weston, Florida 33326
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Revised: August 2018

## **PRINCIPAL DISPLAY PANEL - 10 mg**

**APOTEX CORP.** NDC 60505-0147-1

**Loratadine Tablets 10 mg**

***Non-Drowsy\****

*Antihistamine/Original Prescription Strength*

**†Compare to the active ingredient in Claritin® Tablets**

**Indoor & Outdoor Allergies**

**24 hour**

**Relief of**

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

- Itchy Throat or Nose



LORATADINE

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60505-0147
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	

Product Characteristics

Color	WHITE	Score	no score
Shape	OVAL	Size	8mm
Flavor		Imprint Code	LOR;10;APO
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60505-0147-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/24/2005	06/30/2022
2	NDC:60505-0147-8	1000 in 1 BOTTLE; Type 0: Not a Combination Product	01/24/2005	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA		ANDA076471	01/24/2005	

**Labeler -** Apotex Corp. (845263701)

Revised: 11/2023

Apotex Corp.