

**NUVICARE-ALLERGY RELIEF LORATADINE TABLETS, 10 MG- loratadine tablet**  
**NUVICARE LLC**

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**Nuvicare-Allergy Relief Loratadine Tablets, 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 respiratory allergies:

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

**Warnings**

**Do not use** if you ever have 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**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84324-027
Route of Administration	oral		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>LORATADINE</b> (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg

**Inactive Ingredients**

Ingredient Name	Strength
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>povidone</b> (UNII: FZ989GH94E)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	8mm
<b>Flavor</b>		<b>Imprint Code</b>	P;10
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84324-027-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2025	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA214684	07/01/2025	

**Labeler** - NUVICARE LLC (119257565)

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NUVICARE LLC