

**LORATADINE- loratadine tablet**  
**Northstar RxLLC**

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**NorthStar RxLLC Loratadine Tablets 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?

**1-800-206-7821**

## Principal Display Panel

Compare to active ingredient of Claritin®

Non-Drowsy\*

Original Prescription Strength

Loratadine Tablets, USP

10 mg

Antihistamine

Indoor & Outdoor Allergies

24 Hour

Relief of:

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

Actual Size

\*When taken as directed.

See Drug Facts Panel.

◆NORTHSTARx®

30 Tablets



## LORATADINE

loratadine tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:16714-898
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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<b>LORATADINE</b> (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)		LORATADINE	10 mg	
<b>Inactive Ingredients</b>				
<b>Ingredient Name</b>			<b>Strength</b>	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)				
<b>Product Characteristics</b>				
<b>Color</b>	WHITE	<b>Score</b>	no score	
<b>Shape</b>	OVAL	<b>Size</b>	8mm	
<b>Flavor</b>		<b>Imprint Code</b>	L612	
<b>Contains</b>				
<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:16714-898-01	1 in 1 CARTON	02/20/2019	
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:16714-898-02	1 in 1 CARTON	02/20/2019	
2		100 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:16714-898-03	1 in 1 CARTON	02/20/2019	
3		300 in 1 BOTTLE; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
ANDA	ANDA076301	02/20/2019		

**Labeler** - Northstar RxLLC (830546433)

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Northstar RxLLC