

**EQUALINE ALLERGY RELIEF- loratadine tablet**  
**United Natural Foods, Inc. dba UNFI**

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**SuperValu Inc. Allergy Relief 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?

**1-855-423-2630**

## Principal Display Panel

compare to Claritin® Tablets active ingredient

allergy relief

loratadine tablets, 10mg (antihistamine)

non-drowsy\*

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.

30 tablets

30 days of relief

actual size

ORIGINAL PRESCRIPTION STRENGTH



## EQUALINE ALLERGY RELIEF

loratadine tablet

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41163-612
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, UNSPECIFIED</b> (UNII: FZ989GH94E)	

Product Characteristics			
<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>			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41163-612-46	10 in 1 CARTON	02/07/2005	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:41163-612-65	1 in 1 CARTON	02/15/2005	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:41163-612-72	1 in 1 CARTON	03/23/2005	
3		60 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:41163-612-75	1 in 1 CARTON	03/21/2005	
4		90 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:41163-612-60	20 in 1 CARTON	02/08/2005	11/11/2011
5		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
6	NDC:41163-612-95	1 in 1 CARTON	08/31/2009	06/17/2012
6		45 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:41163-612-58	1 in 1 CARTON	10/03/2019	12/31/2021
7		40 in 1 BOTTLE; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076301	02/07/2005	

**Labeler** - United Natural Foods, Inc. dba UNFI (943556183)

Revised: 7/2023

United Natural Foods, Inc. dba UNFI