

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

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

EQUALINE®

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

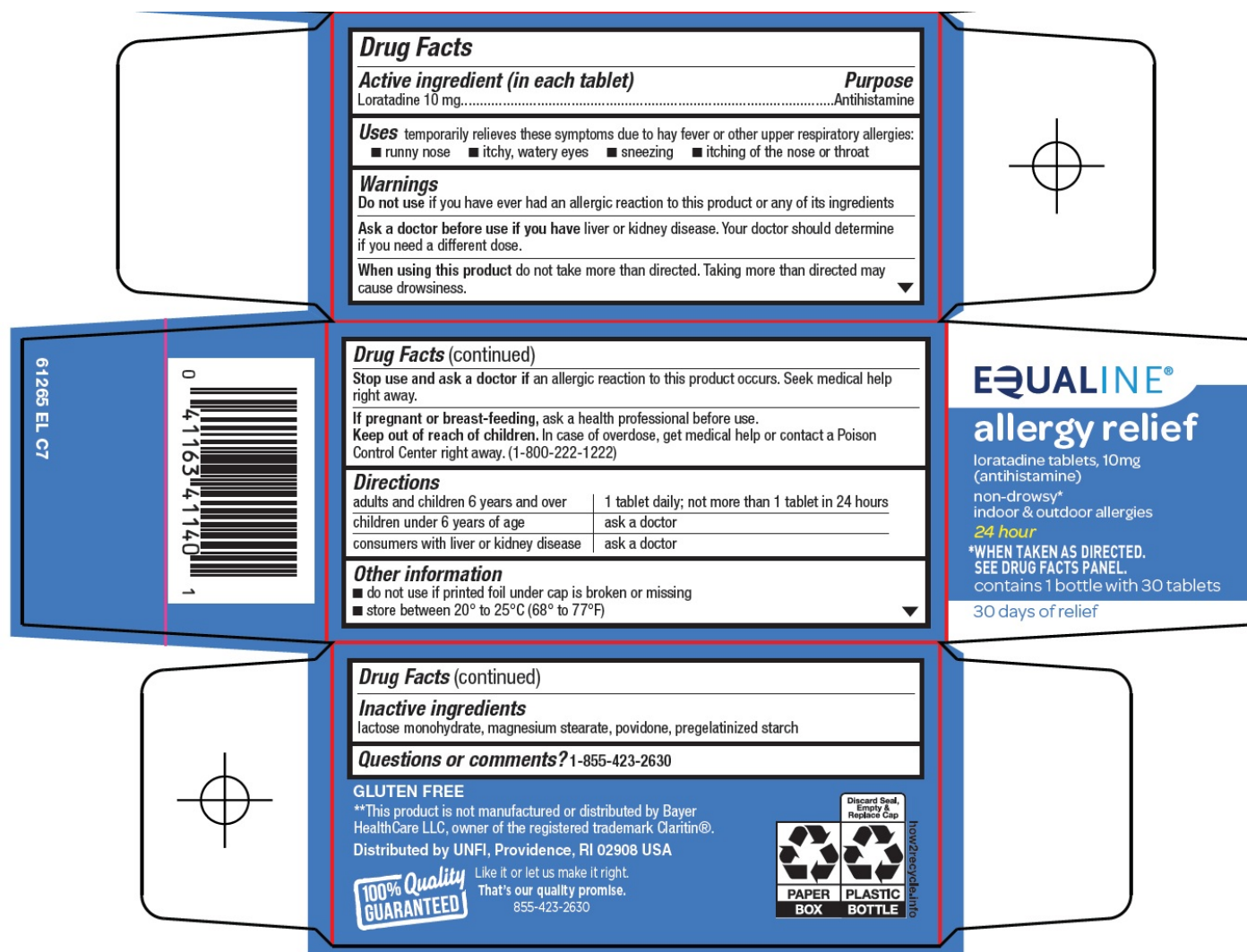
*WHEN TAKEN AS DIRECTED. SEE DRUG FACTS PANEL.

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
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Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	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: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		

8	NDC:41163-612-78	1 in 1 CARTON	12/10/2024	
8		100 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: 12/2024

United Natural Foods, Inc. dba UNFI