

ALLERGY RELIEF- loratadine tablet
Marc Glassman, Inc.

414-Allergy Relief

Active ingredient(s)

Loratadine 10mg

Purpose

Antihistamine

Use(s)

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 accidental overdose, contact a doctor or Poison Control Center (1-800-222-1222) 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

- do not use if blister unit is torn or open

Storage

- store between 20° to 25°C (68° to 77°F)
- protect from excessive moisture

Inactive ingredients

lactose monohydrate, magnesium stearate, pregelatinized starch, sodium starch glycolate

Principal Display Panel

C10-AC1619

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. **You may report side effects to 1-888-952-0050.**
If pregnant or breast-feeding, ask a health professional before use.
Keep out of reach of children. In case of accidental overdose, contact a doctor or Poison Control Center (1-800-222-1222) right away.

Directions

Adults and children 6 years and over | 1 tablet daily: not more than 1 tablet in 24 hours
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Inactive ingredients lactose monohydrate, magnesium stearate, pregelatinized starch, sodium starch glycolate

†This product is not manufactured or distributed by Bayer Healthcare LLC, owner of the registered trademark Claritin®



Distributed By:
Marc Glassman, Inc.
West 130th Street
Cleveland, OH 44130
L82054
R0522

2 25/32

3/4

Marc's

Allergy Relief

30 TABLETS - 10 MG EACH

For Relief of:
24 Hour
• Sneezing
• Itchy, Watery Eyes
• Runny Nose
• Itchy Throat

Actual Size



ANTHISTAMINE

Loratadine

Allergy Relief

Non-Drowsy

†Compare to the active ingredient in Claritin®

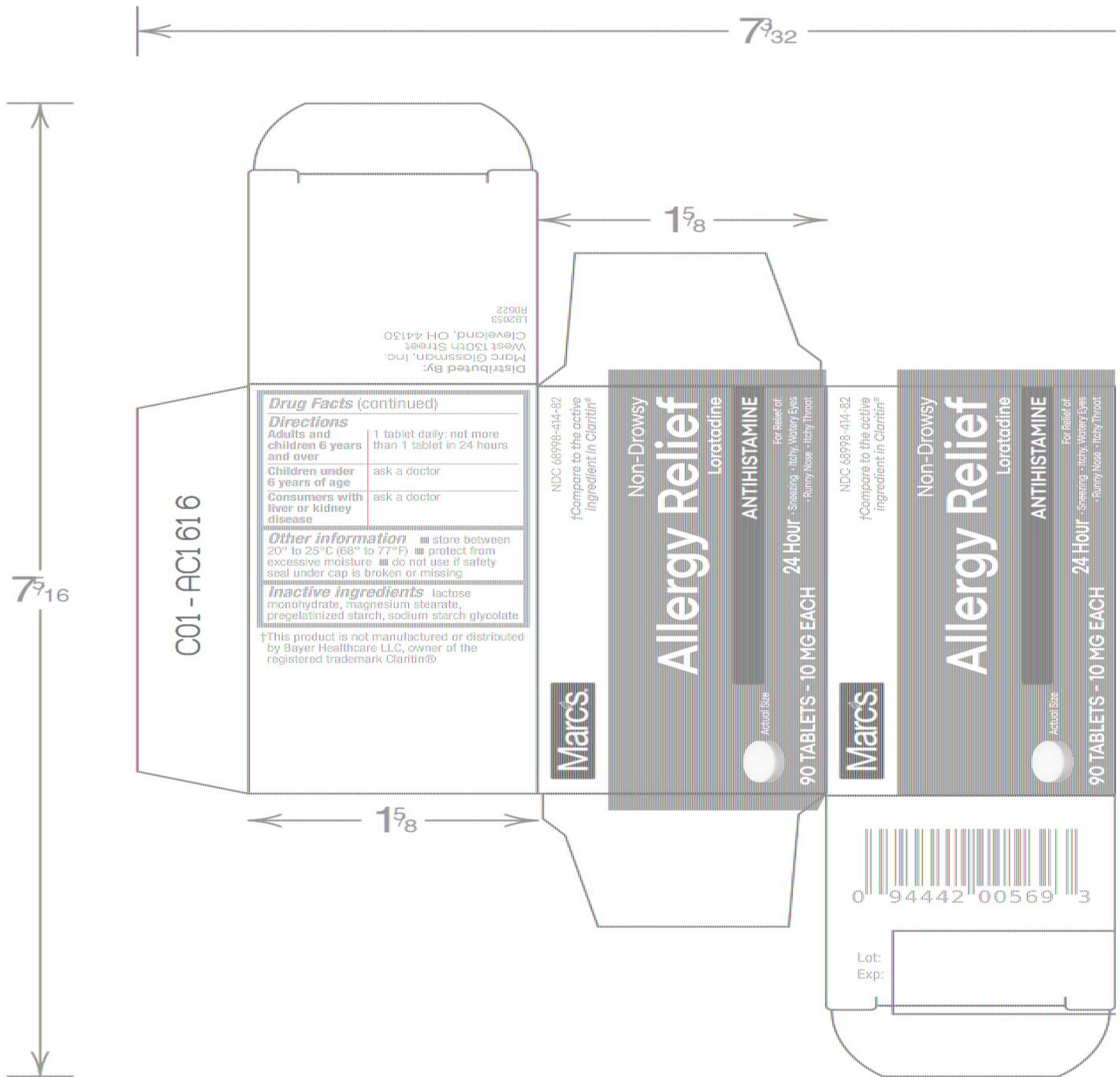
NDC 68998-414-85

Marc's

Allergy Relief

Marc's

Allergy Relief 30



Allergy Relief

ALLERGY RELIEF

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68998-414
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg
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Inactive Ingredients

Ingredient Name	Strength
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM STARCH GLYCOLATE TYPE A (UNII: H8AV0SQX4D)	

Other Ingredients

Ingredient Kind	Ingredient Name	Quantity
May contain	MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
May contain	STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	G;10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68998-414-85	30 in 1 CARTON; Type 0: Not a Combination Product	05/01/2020	
2	NDC:68998-414-82	90 in 1 BOTTLE; Type 0: Not a Combination Product	08/16/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210722	05/01/2020	

Labeler - Marc Glassman, Inc. (094487477)