LORATADINE- loratadine tablet
SAFEWAY INC.

Drug Facts

ACTIVE INGREDIENT (IN EACH TABLET)

Loratadine, USP 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

- store between 20° to 25° C (68° to 77° F)
- protect from excessive moisture
- **TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL IS BROKEN OR MISSING FROM BOTTLE.**

INACTIVE INGREDIENTS

Corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

QUESTIONS OR COMMENTS?

Call **1-888-732-3929**

PRINCIPAL DISPLAY PANEL - 10 mg Tablet Bottle Carton

NDC 21130-526-38

Signature
care™
Quality Guaranteed

24 HOUR | ORIGINAL PRESCRIPTION STRENGTH

Allergy Relief

Loratadine Tablets, USP 10 mg
Antihistamine

Compare to
Claritin®
active ingredient†

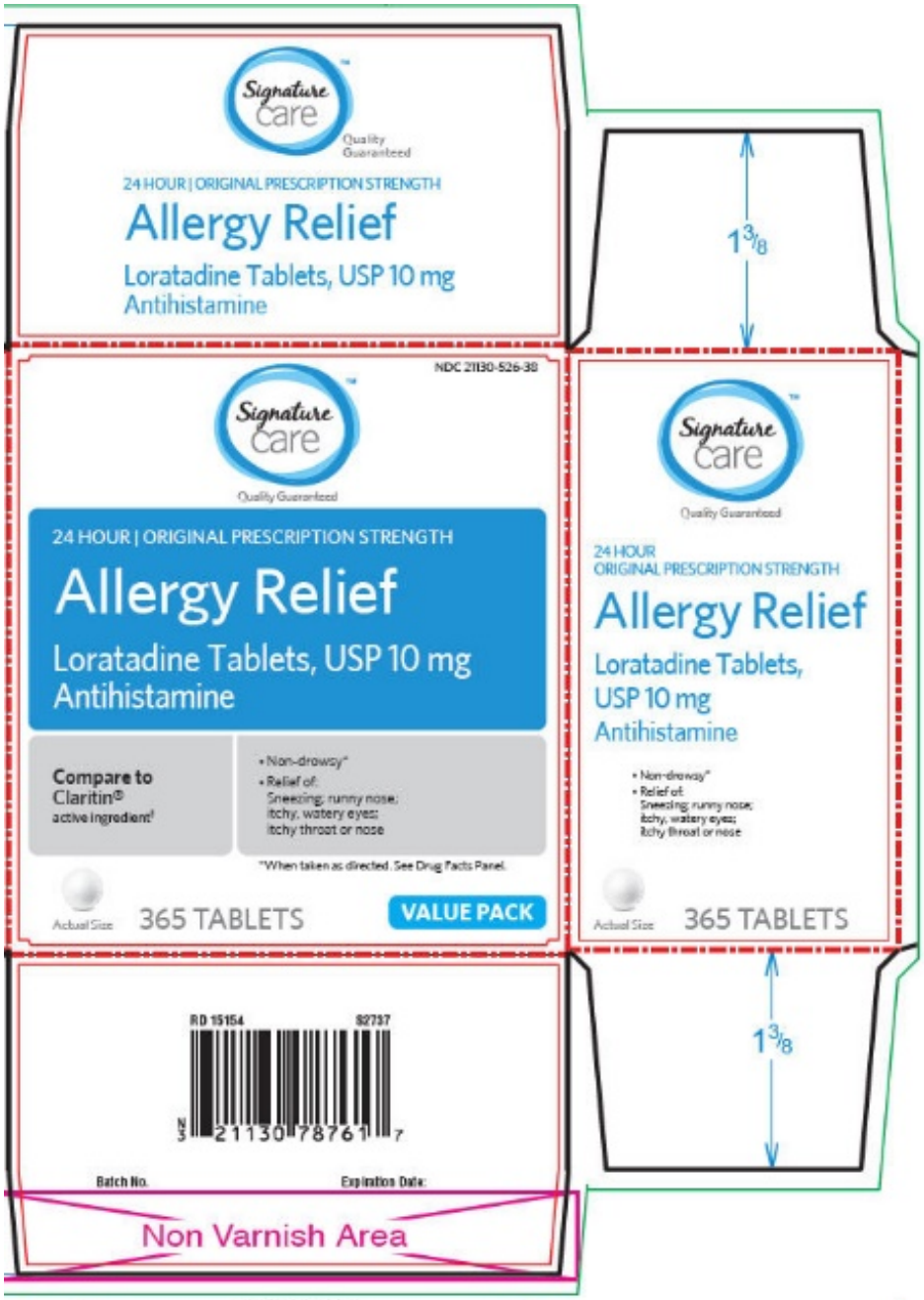
- Non-drowsy*
- Relief of:
  Sneezing; runny nose;
  itchy, watery eyes;
  itchy throat or nose

*When taken as directed. See Drug Facts Panel.

Actual Size

365 TABLETS

VALUE PACK


Signature care
Quality Guaranteed
24 HOUR | ORIGINAL PRESCRIPTION STRENGTH
Allergy Relief
Loratadine Tablets, USP 10 mg
Antihistamine
1 3/8
NDC 21130-526-38
Signature care
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Loratadine Tablets, USP 10 mg
Antihistamine
Compare to Claritin®
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Sneezing; runny nose;
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*When taken as directed. See Drug Facts Panel.
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Antihistamine
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• Relief of:
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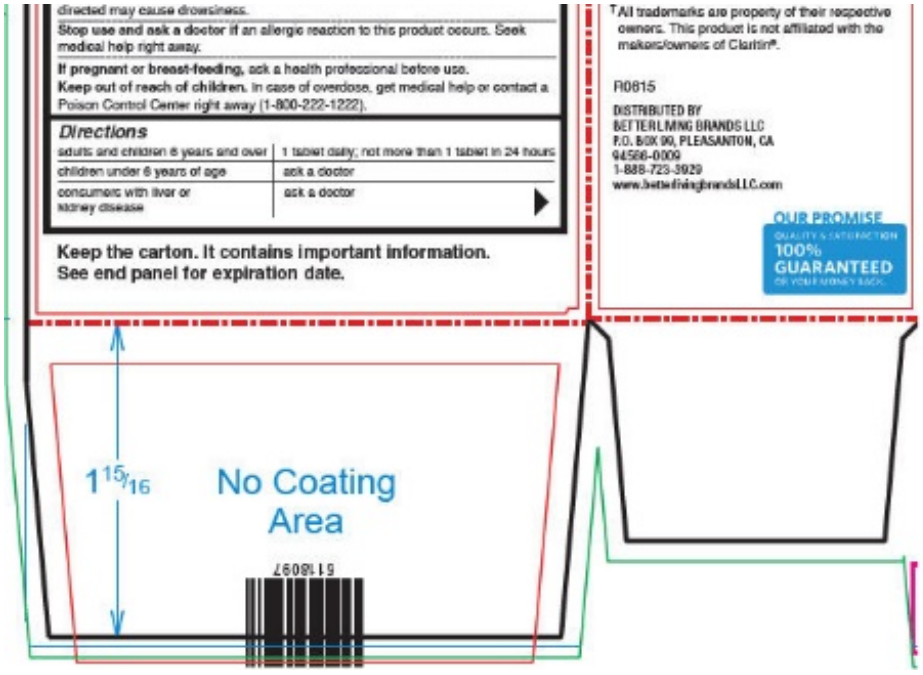
Actual Size
365 TABLETS
1 3/8
RD 15154
82737
21130 78761 7
Batch No.
Expiration Date:
Non Varnish Area
5118097
No Coating Area
1 15/16
Drug Facts
Active Ingredient (in each tablet)
Purpose
Loratadine, USP 10 mg
Antihistamine
Uses temporarily relieves these symptoms due to hay fever or other upper respiratory 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
Drug Facts (continued)
Other Information
■ store between 20° to 25° C (68° to 77° F)
■ protect from excessive moisture
■ TAMPER EVIDENT: DO NOT USE IF IMPRINTED SEAL IS BROKEN OR MISSING FROM BOTTLE.
Inactive Ingredients corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch
Questions? call 1-888-723-3929



LORATADINE

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:21130-526
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients	
Ingredient Name	**Strength**
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics			
Color	white (White to Off-White)	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	RX526
Contains			

Packaging				
			Marketing Start	**Marketing End**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-526-69	1 in 1 CARTON	06/06/2009	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:21130-526-31	3 in 1 CARTON	06/06/2009	
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:21130-526-43	1 in 1 CARTON	06/06/2009	
3		45 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:21130-526-13	1 in 1 CARTON	06/06/2009	
4		120 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:21130-526-38	1 in 1 CARTON	06/06/2009	
5		365 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	ANDA076134	06/06/2009	

Labeler - SAFEWAY INC. (009137209)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	MANUFACTURE(21130-526)

Revised: 12/2021 SAFEWAY INC.