TOPCARE ALLERGY RELIEF 24 HOUR- loratadine tablet Topco Associates LLC

Topco Associates LLC. 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	1 tablet daily; not more than 1 tablet in 24 hours
over	
children under 6 years of age	ask a doctor
consumers with liver or kidney	ask a doctor
disease	

Other information

- do not use if blister unit is broken or torn
- store between 20° to 25°C (68° to 77°F)
- protect from excessive moisture

Inactive ingredients

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

Questions or comments?

1-888-423-0139

Principal Display Panel

COMPARE TO CLARITIN® TABLETS ACTIVE INGREDIENT

NON-DROWSY*

ORIGINAL PRESCRIPTION STRENGTH

Allergy Relief

LORATADINE TABLETS, 10 mg • ANTIHISTAMINE

INDOOR & OUTDOOR ALLERGIES

24 HOUR

RELIEF OF:

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

10 TABLETS

*When taken as directed. See Drug Facts Panel.

actual size



This TopCare® product is laboratory tested to guarantee its highest quality. Your total satisfaction is guaranteed.

†This product is not manufactured or distributed by Bayer HealthCare LLC, distributor of Claritin® Tablets.



String Facts

call 1-888-423-0139 Scan here for more

topcare@topco.com www.topcarebrand.com @10PC0 PERAGE21 QUESTIONS ? 1-888-423-0139 ELK GROVE VILLAGE, IL 600 07 DISTRIBUTED BY TOPOD ASSOCIATES LLC



Questions or comments?1 488-423-0139	if pregnant or bre set-feeding, ask a heath professional before use. ▶
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*When taken as directed. See Drug Facts Panel.

NDC 36800-672-46

10 TABLETS FOR 10 DAYS OF RELIEF NON-DROWSY*

ORIGINAL PRESCRIPTION STRENGTH

LORATADINE TABLETS, 10 mg · ANTIHISTAMINE

INDOOR & OUTDOOR ALLERGIES



COMPARE TO CLARITIN® TABLETS ACTIVE INGREDIENT[†]

NON-DROWSY*

ORIGINAL PRESCRIPTION STRENGTH

Drug Facts (continued)

rgy Relief

LORATADINE TABLETS, 10 mg • ANTIHISTAMINE

INDOOR & OUTDOOR ALLERGIES



RELIEF OF:

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

10 TABLETS

*When taken as directed. See Drug Facts Panel.

actual size



TOPCARE ALLERGY RELIEF 24 HOUR

loratadine tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:36800-612

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AIO3BO7ON) (LORATADINE - UNII:7AIO3BO7ON)	LORATADINE	10 ma

Inactive Ingredients			
Ingredient Name	Strength		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
POVIDONE, UNSPECIFIED (UNII: FZ 989GH94E)			

Product Characteristics				
Color	WHITE	Score	no score	
Shape	OVAL	Size	8mm	
Flavor		Imprint Code	L612	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:36800-612- 65	1 in 1 CARTON	01/25/2005		
1		30 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:36800-612- 72	1 in 1 CARTON	03/24/2005	08/17/2011	
2		60 in 1 BOTTLE; Type 0: Not a Combination Product			
3	NDC:36800-612- 76	1 in 1 CARTON	12/18/2009	11/20/2021	
3		120 in 1 BOTTLE; Type 0: Not a Combination Product			
4	NDC:36800-612- 87	1 in 1 CARTON	12/18/2009		
4		300 in 1 BOTTLE; Type 0: Not a Combination Product			
	NDC-20000 612				

5	NDC:30800-012-	10 in 1 CARTON	01/28/2005	
5		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
6	NDC:36800-612- 58	1 in 1 CARTON	02/19/2014	
6		40 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:36800-612- 03	1 in 1 CARTON	03/05/2014	
7		70 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information			
			Marketing End Date
ANDA	ANDA076301	01/25/2005	

Labeler - Topco Associates LLC (006935977)

Revised: 11/2021 Topco Associates LLC