CLARITIN LIQUI-GELS- loratadine capsule, liquid filled
Bayer HealthCare LLC.

Claritin ®

Liqui-Gels ®

Drug Facts

Active ingredient (in each capsule)

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.

Directions

adults and children 6	1 capsule daily; not more than 1

years and over	capsule in 24 hours
children under 6 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- safety sealed: do not use if the individual blister unit imprinted with Claritin® Liqui-Gels® is open or torn
- store between 20° to 25°C (68° to 77°F)
- protect from freezing

Inactive ingredients

caprylic/capric glycerides, FD&C blue no.1, gelatin, glycerin, pharmaceutical ink, polysorbate 80, povidone, purified water, sorbitol

Questions or comments?

1-800-CLARITIN (1-800-252-7484)

Pat.: patents.livewell.bayer.com

Whippany, NJ 07981

PRINCIPAL DISPLAY PANEL - 30 Capsule Carton

Non-Drowsy*

Claritin ®

Liqui-Gels®

24

Hour

Relief of:

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

loratadine 10 mg/antihistamine

Indoor & Outdoor Allergies

Allergies

***When taken as directed.**

See Drug Facts Panel.

30 LIQUID-FILLED

CAPSULES

30 Count DFL

CLARITIN LIQUI-GELS

loratadine capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-7200
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**
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE (UNII: FZ989GH94E)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

Product Characteristics			
Color	blue (Clear Blue)	**Score**	no score
Shape	CAPSULE	**Size**	10mm
Flavor		**Imprint Code**	C;10
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11523-7200-1	1 in 1 CARTON	06/16/2008	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11523-7200-2	3 in 1 CARTON	06/16/2008	01/01/2016
2		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:11523-7200-3	4 in 1 CARTON	06/16/2008	01/01/2016
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:11523-7200-5	1 in 1 POUCH; Type 0: Not a Combination Product	06/16/2008	01/01/2016
5	NDC:11523-7200-4	7 in 1 CARTON	06/16/2008	01/01/2016
5		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
6	NDC:11523-7200-6	3 in 1 CARTON	06/16/2008	
6		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
7	NDC:11523-7200-7	5 in 1 CARTON	06/16/2008	01/01/2016
7		9 in 1 BLISTER PACK; Type 0: Not a Combination Product		
8	NDC:11523-7200-8	4 in 1 CARTON	06/16/2008	
8		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
9	NDC:11523-7200-9	6 in 1 CARTON	06/16/2008	
9		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA021952	06/16/2008	

CLARITIN LIQUI-GELS

loratadine capsule, liquid filled

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11523-7333
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
GLYCERIN (UNII: PDC6A3C0OX)	
POVIDONE (UNII: FZ989GH94E)	
WATER (UNII: 059QF0KO0R)	
SORBITOL (UNII: 506T60A25R)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
GELATIN (UNII: 2G86QN327L)	

Product Characteristics

Color	blue (Clear Blue)	**Score**	no score
Shape	CAPSULE	**Size**	10mm
Flavor		**Imprint Code**	C;10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-7333-1	1 in 1 CARTON	06/16/2008	
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11523-7333-3	4 in 1 CARTON	06/16/2008	
2		9 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:11523-7333-4	7 in 1 CARTON	06/16/2008	
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:11523-7333-5	10 in 1 CARTON	06/16/2008	
4		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA021952	06/16/2008	

Labeler - Bayer HealthCare LLC. (112117283)

Revised: 2/2026 Bayer HealthCare LLC.