

CLARITIN REDITABS- loratadine tablet, orally disintegrating
Bayer Healthcare LLC.

Claritin®

RediTabs®

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.

Directions

- place 1 tablet on tongue; tablet disintegrates, with or without water

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

- safety sealed: do not use if the individual blister unit imprinted with Claritin® RediTabs® is open or torn
- store between 20° to 25°C (68° to 77°F)
- use tablet immediately after opening individual blister
- complies with USP Procedure 2 for Assay and Organic Impurities and Test 2 for Disintegration

Inactive ingredients

anhydrous citric acid, gelatin, mannitol, mint flavor

Questions or comments?

1-800-CLARITIN (1-800-252-7484) or www.claritin.com

**Distributed by MSD Consumer Care, Inc.,
PO Box 377, Memphis, TN 38151 USA,
a subsidiary of Merck & Co., Inc., Whitehouse Station, NJ USA.**

PRINCIPAL DISPLAY PANEL - 10 Tablet Carton

Original Prescription Strength

NDC 11523-7157-2

Non-Drowsy*

Claritin®

RediTabs®

loratadine 10 mg/antihistamine

Indoor & Outdoor

Allergies

****When taken as directed. See Drug Facts Panel.***

24

Hour

Relief of:

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

- *Itchy Throat
or Nose*

**No Water Needed
Melts in Your Mouth**

10

ORALLY

DISINTEGRATING TABLETS

Claritin® Reditabs®

TABLETS

24
Hour

10 ORALLY DISINTEGRATING TABLETS
FOR 10 DAYS OF RELIEF

Original Prescription Strength

NDC 11523-7157-2

Non-Drowsy*

Claritin®

Reditabs®

loratadine 10 mg/antihistamine

Indoor & Outdoor Allergies

24
Hour

Relief of:

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

No Water Needed
Melts in Your Mouth



10 ORALLY DISINTEGRATING TABLETS

*When taken as directed. See Drug Facts Panel.

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3. Place the tablet on tongue and close mouth. The tablet will disintegrate.



2. Gently push tablet out.



1. Peel back outer edge.



Follow these directions carefully. Do not attempt to push the tablet through the foil.

24
Hour

- No Water Needed
- Melts in your mouth

Claritin® Reditabs®

1-800-CLARITIN (1-800-252-7484) or www.claritin.com

Questions or comments?

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

Place 1 tablet on tongue; tablet disintegrates, with or without water.

adults and children 6 years and over
1 tablet daily; not more than 1 tablet in 24 hours

ask a doctor

children under 6 years of age
ask a doctor

consumers with liver or kidney disease
ask a doctor

Drug Facts

Active ingredient (in each tablet)

Loratadine 10 mg.....Antihistamine

Purpose

Uses temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- sneezing
- runny nose
- itchy, watery eyes
- 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.

Drug Facts (continued)

Other information

- safety sealed: do not use if the individual blister unit imprinted with Claritin® RediTabs® is open or torn
- store between 20° to 25° C (68° to 77° F)
- use tablet immediately after opening individual blister
- complies with USP Procedure 2 for Assay and Organic Impurities and Test 2 for Disintegration

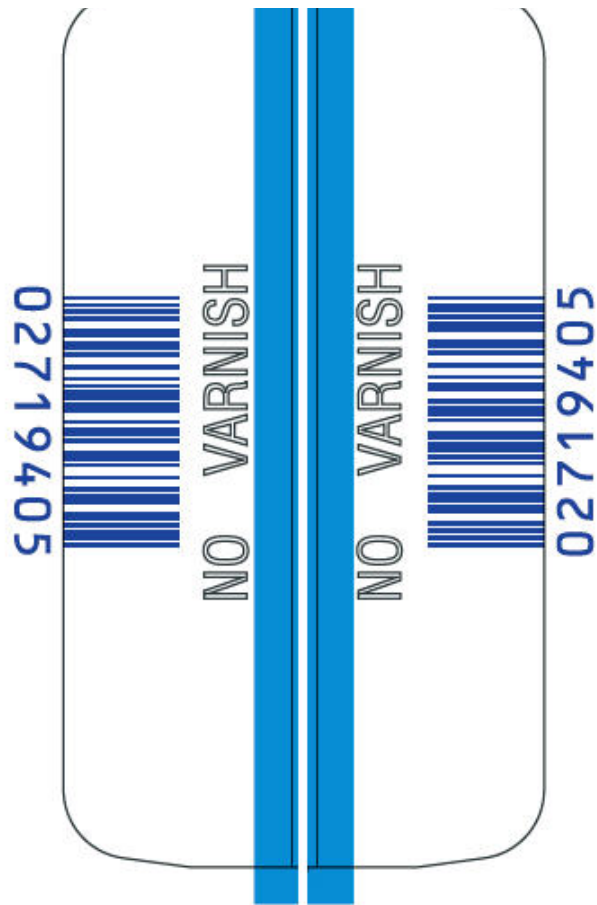
Inactive ingredients

anhydrous citric acid, gelatin, mannitol, mint flavor

NO VARNISH



NO VARNISH
CODE AREA



PRINCIPAL DISPLAY PANEL - 60 Tablet Twin Pack Carton

TWIN PACK

**Two 30ct Cartons
60 RediTabs[®]**

Original Prescription Strength

NDC 11523-4329-1

Non-Drowsy*

Claritin[®]

RediTabs[®]

loratadine 10 mg/antihistamine

**Indoor & Outdoor
Allergies**

***When taken as directed. See Drug Facts Panel.**

**24
Hour**

Relief of:

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

or Nose

**No Water Needed
Melts in Your Mouth**

**60
ORALLY
DISINTEGRATING TABLETS**

TWIN PACK

Two 30ct Cartons 60 Reditabs[®]

Original Prescription Strength

NDC 11523-4329-1

Non-Drowsy*
Claritin[®]

**24
Hour**

No Water Needed
Melts in Your Mouth

Reditabs[®]

loratadine 10 mg/antihistamine

Relief of:

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

Indoor & Outdoor
Allergies

60 ORALLY
DISINTEGRATING TABLETS

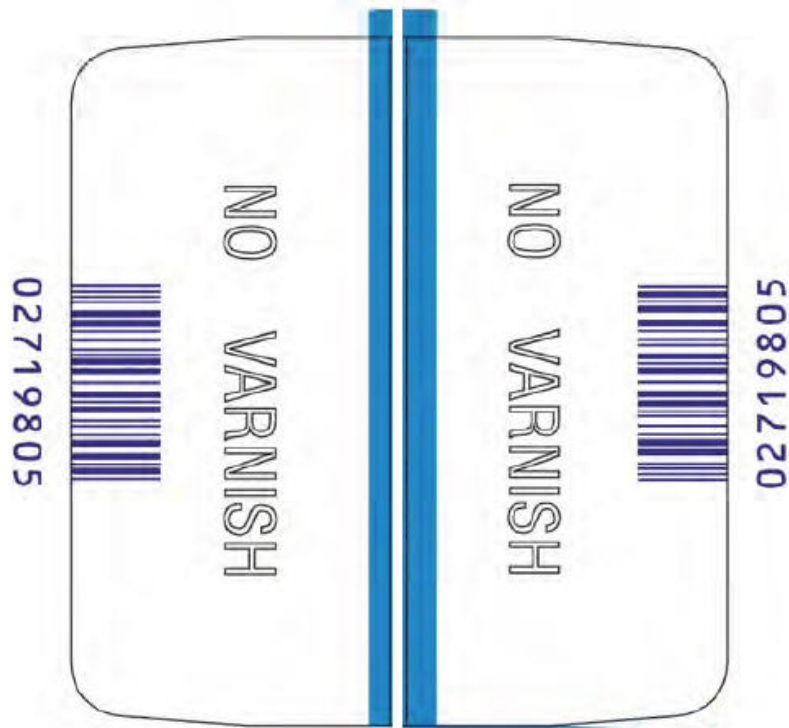
*When taken as directed. See Drug Facts Panel.

32437-00

NO

VARNISH

Drug Facts		Drug Facts (continued)	
Active ingredient (in each tablet) Loratadine 10 mg.....		Other information	
Purpose Antihistamine		■ safety sealed: do not use if the individual blister unit imprinted with Claritin [®] Reditabs [®] is open or torn	
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		■ store between 20° to 25° C (68° to 77° F)	
Warnings		■ use tablet immediately after opening individual blister	
Do not use if you have ever had an allergic reaction to this product or any of its ingredients.		■ complies with USP Procedure 2 for Assay and Organic Impurities and Test 2 for Disintegration	
Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you need a different dose.		Inactive ingredients anhydrous citric acid, gelatin, mannitol, mint flavor	
When using this product do not take more than directed. Taking more than directed may cause drowsiness.		Questions or comments? 1-800-CLARITIN (1-800-252-7484) or www.claritin.com	
Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away.		Claritin[®] Reditabs[®] • No Water Needed • Melts in your mouth 24 Hour	
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			
■ place 1 tablet on tongue; tablet disintegrates, with or without water			
adults and children 6 years and over	1 tablet daily; not more than 1 tablet in 24 hours	Follow these directions carefully. Do not attempt to push the tablet through the foil.	
children under 6 years of age	ask a doctor		
consumers with liver or kidney disease	ask a doctor		
			
		1. Peel back outer edge.	
		2. Gently push tablet out.	
		3. Place the tablet on tongue and close mouth. The tablet will disintegrate.	



CLARITIN REDITABS

loratadine tablet, orally disintegrating

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11523-7157
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
GELATIN (UNII: 2G86QN327L)	
MANNITOL (UNII: 3OWL53L36A)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	12mm
Flavor	MINT	Imprint Code	C10
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-7157-2	1 in 1 CARTON	11/27/2002	
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11523-7157-3	3 in 1 CARTON	11/27/2002	
2		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:11523-7157-4	2 in 1 CARTON	11/27/2002	
3		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:11523-7157-7	4 in 1 CARTON	11/27/2002	
4		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:11523-7157-8	5 in 1 CARTON	11/27/2002	
5		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
6	NDC:11523-7157-9	7 in 1 CARTON	11/27/2002	
6		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	NDA020704	11/27/2002	

CLARITIN REDITABS			
loratadine tablet, orally disintegrating			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11523-4329
Route of Administration	ORAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)		LORATADINE	10 mg
Inactive Ingredients			
Ingredient Name			Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)			

GELATIN (UNII: 2G86QN327L)	
MANNITOL (UNII: 3OWL53L36A)	

Product Characteristics

Color	white	Score	no score
Shape	ROUND	Size	12mm
Flavor	MINT	Imprint Code	C10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11523-4329-1	2 in 1 PACKAGE, COMBINATION	11/27/2002	
1		3 in 1 CARTON		
1		10 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11523-4329-2	6 in 1 CARTON	11/27/2002	
2		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	NDA020704	11/27/2002	

Labeler - Bayer Healthcare LLC. (112117283)

Revised: 11/2023

Bayer Healthcare LLC.