

**LORATADINE ANTIHISTAMINE- loratadine tablet**  
**Unit Dose Services**

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**Perrigo Loratadine Tablets, 10 mg 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 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 printed foil under cap is broken or missing
- store between 20° to 25°C (68° to 77°F)

## Inactive ingredients

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

## Questions or comments?

1-800-719-9260

## HOW SUPPLIED

Product: 50436-3502

NDC: 50436-3502-1 30 TABLET in a BOTTLE

## LORATADINE ANTIHISTAMINE (LORATADINE) TABLET

NDC: 50436-3502-1  
**LORATADINE 10MG 30 TABLETS**  
30 TABLETS, EACH CONTAINING  
10MG LORATADINE  
LOT: XXXXX EXP: XXXXXX  
DIST BY: PERRIGO  
ALLEGAN, MI 49010  
WARNING: KEEP OUT OF REACH OF CHILDREN  
STORE AT 20°-25°C (68°-77°F)  
SEE PACKAGE INSERT FOR DOSAGE INFORMATION  
MFG NDC: 45802-650-87  
MFG LOT: XXXXX  
PKG BY: UNIT DOSE SERVICES LLC, DANIA, FL



## LORATADINE ANTIHISTAMINE

loratadine tablet

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50436-3502(NDC:45802-650)
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
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LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50436-3502-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	10/15/2008	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076301	10/15/2008	

**Labeler** - Unit Dose Services (831995316)

Establishment			
Name	Address	ID/FEI	Business Operations
Unit Dose Services		831995316	REPACK(50436-3502) , RELABEL(50436-3502)

Revised: 7/2017

Unit Dose Services