LORATADINE- loratadine tablet
Major Pharmaceuticals

Loratadine tablets

Active ingredient
Loratidine USP 10 mg

Purpose
Antihistamine

Use
Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies, runny nose, itchy, watery eyes, sneezing, itching of the nose and throat.

Warning
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. You doctor should determine if you need a different dose.

Ask a doctor or pharmacist before use if you are

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.

Pregnancy/breast-feeding warning
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 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 at 20º - 25ºC (68º -77ºF); excursions permitted to 15º - 30º C (59º - 86º F). [See USP Controlled Room Temperature]
Protect from excessive moisture.

Inactive ingredients
Corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch.

Questions?
Questions or comments? (800) 616-2471
Distributed by:
MAJOR® PHARMACEUTICALS
31778 Enterprise Drive, Livonia, MI 48150 USA
Re-Order No. 301592

PRINCIPAL DISPLAY PANEL LORATADINE TABLETS, USP 10MG

PHARMACEUTICALS

NDC 0904-6074-61
Unit Dose
LORATADINE TABLETS, USP
10 mg.
ANTIHISTAMINE

Relief of: Sneezing, Runny Nose, Itchy, Watery Eyes, Itchy Throat or Nose

100 TABLETS

MAJOR®
PHARMACEUTICALS

NDC 0904-6074-61
Unit Dose
LORATADINE 10mg Tablets

Drug Facts

Active ingredient (in each tablet)	***Purpose***
Loratadine USP, 10mg	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 and 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 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 at 20°-25°C (68°-77°F); excursions permitted to 15°-30°C (59°-86°F). [See USP Controlled Room Temperature] ■ protect from excessive moisture.

Inactive ingredients: Corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch.

Questions or comments? (800) 616-2471

LORATADINE

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG LABEL	**Item Code (Source)**	NDC:0904-6074(NDC:51660-526)
Route of Administration	ORAL	**DEA Schedule**	

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LORATADINE (LORATADINE)	LORATADINE	10 mg

Inactive Ingredients	
Ingredient Name	**Strength**
STARCH, CORN	
LACTOSE MONOHYDRATE	
MAGNESIUM STEARATE	

Product Characteristics			
Color	WHITE	**Score**	no score
Shape	ROUND	**Size**	6mm
Flavor		**Imprint Code**	RX526
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:0904-6074-61	100 in 1 BOX, UNIT-DOSE; Combination Product Type = C112160	06/15/2010	08/31/2017

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
ANDA	ANDA076134	06/15/2010	

Labeler - Major Pharmaceuticals (191427277)

Revised: 11/2016

Major Pharmaceuticals