

LORATADINE- loratadine tablet
ST. MARY'S MEDICAL PARK PHARMACY

Loratadine Tablets, 10 mg

ACTIVE INGREDIENT(S)

Loratadine 10 mg

PURPOSE

Antihistamine

USE(S)

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

- Tamper-evident: do not use if foil seal under cap, printed with "SEALED for YOUR PROTECTION" is missing, open or broken

STORAGE

- store between 20° to 25°C (68° to 77°F)

INACTIVE INGREDIENTS

Lactose monohydrate, magnesium stearate, sodium starch glycolate and starch maize pregelatinized

QUESTIONS OR COMMENTS

Contact 1-888-588-1418 Mon-Fri 8:00 AM EST to 5:00 PM EST

PRINCIPAL DISPLAY PANEL

NDC 60760-609-00

Loratadine
 Antihistamine
 Allergy Relief
 Tablets, USP
10mg

QTY: 100
 LOT# XXXXXXX
 EXP XX-XX
 RX# ???????

MANUFACTURED BY:
 Camber Consumer Care Inc.
 Piscataway, NJ 08854, USA

60760-589-00



Loratadine
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 Tablets, USP
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 **PACKAGED BY:**
 St. Mary's
 10860 MAVINEE DR.
 ORO VALLEY, AZ 85737
 MANAGED PHARMACY PROGRAMS

TAKE AS DIRECTED



Rx only STORE AT CONTROLLED ROOM TEMPERATURE 15°- 30° C (59°-86°F)

LORATADINE

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:60760-609(NDC:69230-328)
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
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STARCH, CORN (UNII: O8232NY3S)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)	

Product Characteristics

Color	white (White to off white)	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	G;10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start	Marketing End
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#	Item Code	Package Description	Date	Date
1	NDC:60760-609-00	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/30/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210722	07/31/2022	

Labeler - ST. MARY'S MEDICAL PARK PHARMACY (063050751)

Establishment			
Name	Address	ID/FEI	Business Operations
ST. MARY'S MEDICAL PARK PHARMACY		063050751	relabel(60760-609) , repack(60760-609)

Revised: 12/2023

ST. MARY'S MEDICAL PARK PHARMACY