

GOOD NEIGHBOR PHARMACY LORATADINE- loratadine tablet
Preferred Pharmaceuticals Inc.

Amerisource Bergen 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 {Bottle Only}
- store between 20° to 25°C (68° to 77°F)
- protect from excessive moisture {Blister Only}

Inactive ingredients

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

Questions or comments?

1-800-719-9260

Repackaged By: Preferred Pharmaceuticals Inc.

Principal Display Panel

Compare to Claritin® Tablets active ingredient

Original Prescription Strength

Loratadine tablets, 10 mg

Antihistamine

Indoor & Outdoor Allergies

24 HOUR

Relief of:

Sneezing

Runny Nose

Itchy, Water Eyes

Itchy Throat or Nose

*When taken as direction.

See Drug Facts Panel.

Non-Drowsy*

actual size

Loratadine
Tablets 10mg

Generic for Claritin

Active ingredient (in each tablet) Loratadine 10mg.....Antihistamine


Pkg Size: Exp Date:

Lot#: Batch#: Ins:

Mfg: Good Neighbor Pharmacy Prod#:

Warning


Store at 20°- 25°C (68°- 77°F). 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. 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. Keep this and all medication out of the reach of children. If pregnant or breast feeding, ask a health professional before use. Tablet is oval, white, and imprinted with L612



Directions English

Use as directed by your doctor

Take ___ tablet(s) every ___ hours.



Instructions Espanol:

Uso según lo dirigido por su doctor

Toma ___ tableta(s) cada ___ horas.

CAUTION: Federal law PROHIBITS transfer of this drug to any person other than the patient for whom it was prescribed

Loratadine Tablets 10mg
Qty: Ins:
Lot#: Bat#:
Prod# (NDC):

Loratadine Tablets 10mg
Qty: Ins:
Lot#: Bat#:
Prod# (NDC):

Loratadine Tablets 10mg
Qty: Ins:
Insurance NDC:
Lot#: Bat#:

Loratadine Tablets 10mg
Qty: Ins:
Lot#: Bat#:
Prod# (NDC):

Log

Chart

Billing

Patient

GOOD NEIGHBOR PHARMACY LORATADINE

loratadine tablet

Product Information				
Product Type		HUMAN OTC DRUG	Item Code (Source)	NDC:68788-8614(NDC:24385-471)
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)				
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:68788-8614-0	10 in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2024	
	NDC:68788-8614-0	10 in 1 BOTTLE; Type 0: Not a Combination Product		

2	NDC:68788-8614-1	14 in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2024	
3	NDC:68788-8614-5	15 in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2024	
4	NDC:68788-8614-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2024	
5	NDC:68788-8614-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076301	03/22/2024	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-8614)

Revised: 3/2024

Preferred Pharmaceuticals Inc.