LORATADINE- loratadine tablet
Preferred Pharmaceuticals Inc.

Loratadine Tablets USP 10mg

Drug Facts

Active ingredient (in each tablet)

Loratadine USP 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 an overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

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

store at 20°C to 25°C (68° to 77°F)

protect from excessive moisture

Inactive ingredients

lactose monohydrate, magnesium stearate, pregelatinized starch (maize), sodium starch glycolate.

Questions or comments?

call **1-855-274-4122**

Manufactured by: **Aurobindo Pharma Limited**

Hyderabad-509 302,

INDIA

For BluePoint Laboratories

MADE IN INDIA

Code: TS/DRUGS/22/2009

Issued: 04/2020

Repackaged By: Preferred Pharmaceuticals Inc.

PACKAGE LABEL- PRINCIPAL DISPLAY PANEL - 10mg

NDC 68788-8628

Non-Drowsy*

Loratadine

Tablets USP 10mg

Antihistamine

24 Hour

Relief of:

Sneezing

Runny Nose

Itchy, Watery Eyes

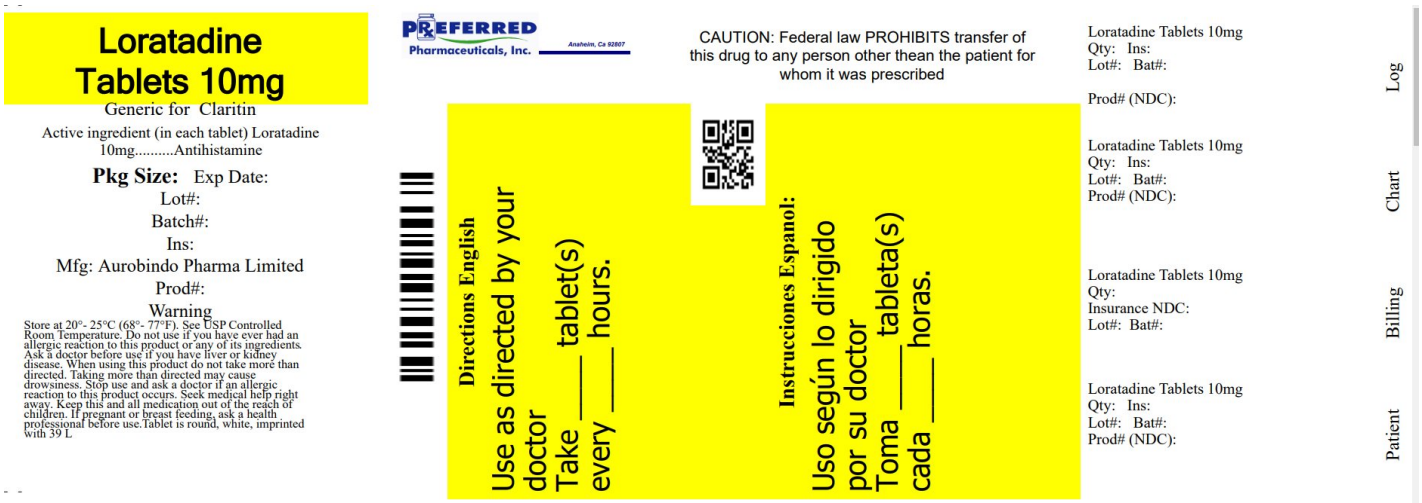
Itchy Throat or Nose

Indoor & Outdoor

Allergies

When taken as directed.

See Drug Facts Panel.



LORATADINE

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68788-8628(NDC:68001-438)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients

Ingredient Name	Strength
MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
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**	39;L
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-8628-0	10 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2024	
2	NDC:68788-8628-1	14 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2024	
3	NDC:68788-8628-5	15 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2024	
4	NDC:68788-8628-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2024	
5	NDC:68788-8628-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/11/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA208314	04/11/2024	

Labeler - Preferred Pharmaceuticals Inc. (791119022)

Registrant - Preferred Pharmaceuticals Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-8628)