LORATADINE- loratadine tablet
DirectRx

Loratadine 10 mg

Antihistamine

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)

- do not use if printed foil under cap is broken or missing
- store between 20° to 25°C (68° to 77°F)

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

1-800-719-9260

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 at 20°-25°C (68°-77°F)
Questions or comments?

1-800-719-9260

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

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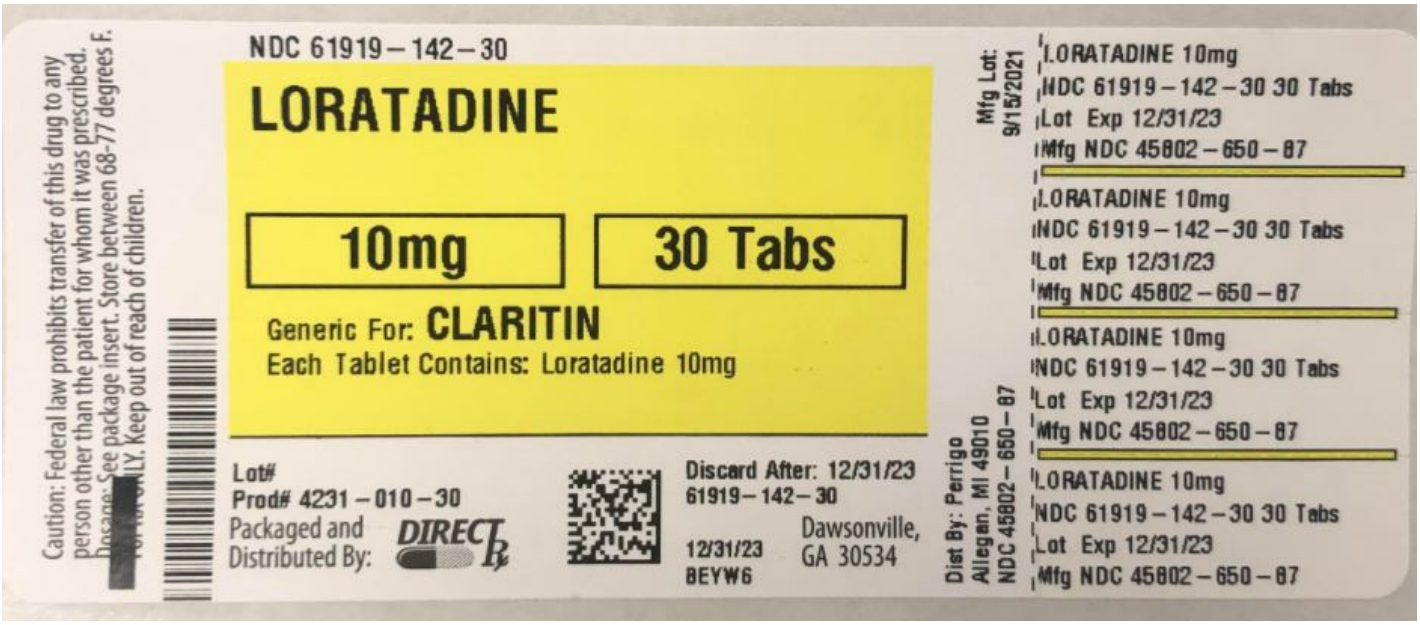
runny nose
•
itchy, watery eyes
•
sneezing
•
itching of the nose or throat



LORATADINE

loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:61919-142(NDC:45802-650)
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)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
POVIDONE (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:61919-142-07	7 in 1 BOTTLE; Type 0: Not a Combination Product	09/13/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076301	09/13/2021	

Labeler - DirectRx (079254320)

Registrant - DirectRx (079254320)

Establishment

Name	Address	ID/FEI	Business Operations
DirectRx		079254320	repack(61919-142)

Revised: 10/2021 DirectRx