# **ALLERGY RELIEF NON DROWSY - loratadine tablet Select Corporation**

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#### **Drug Facts**

Active Ingredients Loratadine USP 10 mg

Antihistamine

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

Uses: • temporarily relieves these symptoms due to hay fever or other upper respiratory allergies: • runny nose • sneezing • itchy, watery eyes • 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.

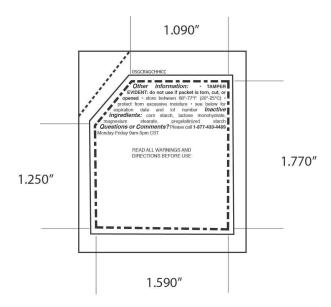
KEEP OUT OF REACH OF CHILDREN.

Inactive

ingredients: corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

MM1





loratadine tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52904-430
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 (UNII: J2B2A4N98G)		
STARCH, CORN (UNII: O8232NY3SJ)		

Product Characteristics			
Color	white (snow white)	Score	no score
Shape	ROUND (RX526)	Size	6 mm
Flavor		Imprint Code	RX526
Contains			

I	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52904-430-01	1 in 1 PACKET		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077153	10/15/2012	

## Labeler - Select Corporation (053805599)

### **Registrant** - Select Corporation (053805599)

Revised: 10/2012 Select Corporation