# UP AND UP ALLERGY RELIEF- loratadine tablet Target Corporation

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#### **Target Corporation Allergy Relief 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
- store between  $20^{\circ}$  to  $25^{\circ}$ C ( $68^{\circ}$  to  $77^{\circ}$ F)

## **Inactive ingredients**

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

Questions? Call 1-888-547-7400

# Principal Display Panel

**CE PACK** Compare to active ingredient in Claritin<sup>®</sup> non-drowsy\* allergy relief loratadine tablets, 10 mg antihistamine 30 days of relief original prescription strength indoor and outdoor allergies 24 hour relief of: sneezing runny nose itchy, watery eyes itchy throat or nose 24 HOUR ACTUAL SIZE **30 TABLETS 30 TABLETS** \*When taken as directed. See drug facts panel.



UP AND UP ALLERGY I	RELIEF						
Product Information							
Product T ype	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-612				
Route of Administration	ORAL						
Active Ingredient/Active Moiety							

Ingredient Name						Basis of St	trength	Strength	
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)						LORATADINE		10 mg	
Ina	active Ingredie	nts							
			Ing	gredient Nam	2			S	trength
LA	<b>CTOSE MONOHYI</b>	DRATE (U	NII: EWQ57Q8	315X)					
MA	GNESIUM STEARA	ATE (UNII:	70097M6I30)	)					
PO	VIDONE (UNII: FZ9	89GH94E	)						
Pro	oduct Characte	ristics							
Col	lor		WHITE	Score			nc	no score 3mm	
Sha	ape		OVAL	Size			8 r		
Fla	vor			Impri	nt Code		Le	512	
Cor	ntains								
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# Marketing InformationMarketing CategoryApplication Number or Monograph CitationMarketing Start DateMarketing End DateANDAANDA7630103/26/201203/26/2012

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