LORATADINE- loratadine tablet Shopko Stores Operating Co., LLC

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

DIRECTIONS

adults and children 6 years and over
children under 6 years of age
consumers with liver or kidney disease ask a doctor1 tablet daily; not more than 1 tablet in 24 hours
ask a doctor

OTHER INFORMATION

- store between 20 and 25° C (68 and 77° F)
- protect from excessive moisture
- TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.

INACTIVE INGREDIENTS

Corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

QUESTIONS?

Call 1-800-406-7984

PRINCIPAL DISPLAY PANEL SHOPKO[®] [†]Compare to the Active Ingredient of Claritin[®] Original Prescription Strength Allergy Relief Loratadine Tablets, USP 10 mg Antihistamine Non-Drowsy^{*} Indoor & Outdoor Allergies 24 Hour Allergy Relief Relief of: Sneezing, Runny Nose, Itchy/Watery Eyes, Itchy Throat or Nose ^{*}When taken as directed. See Drug Facts Panel. Manufactured by: Ohm Laboratories Inc. 5093684/R0412



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Product Informat	lion		_				
Product Type		HUMAN OTC DRUG	Ite m	Code (Sourc	e)	NDC:370	12-526
Route of Administra	tion	ORAL					
Active Ingredient	/Active Moi	e ty					
Ingredient Name					Basis of Strength Strength		
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Inactive Ingredie	nts	T 11 . N					1
CTADOL CODN (UNIT	Ingredient Name						Strength
STARCH, CORN (UNII	: 08232NY3SJ)						
LACTOSE MONOINI	DATE /INTE						
LACTOSE MONOHYI MAGNESIUM STEARA STARCH, PREGELATI	TE (UNII: 7009	7M6I30)					
MAGNESIUM STEARA STARCH, PREGELATI	ATE (UNII: 7009	7M6I30)					
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MAGNESIUM STEARA STARCH, PREGELATI Product Characte Color Shape Flavor	TE (UNII: 7009 INIZED CORN (Pristics white (White to	7M6I30) UNII: 08232NY3SJ)		Size	e	6 m m	1
MAGNESIUM STEARA STARCH, PREGELATI Product Characte Color Shape Flavor Contains Packaging	TE (UNII: 7009 NIZED CORN (ristics white (White to ROUND	7M6I30) UNII: 08232NY3SJ)	Market	Size		6 m m	1 26
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MAGNESIUM STEARA STARCH, PREGELATI Product Characte Color Shape Flavor Contains Packaging # Item Code	TE (UNII: 7009 INIZED CORN (Pristics white (White to ROUND ROUND 10 in 1 Bl	7M6I30) UNII: O8232NY3SJ) off-White) kage Description	Market	Size Imprint Cod		6 mm RX5	1 26
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Labeler - Shopko Stores Operating Co., LLC (023252638)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment								
Name	Address	ID/FEI	Business Operations					
Ohm Laboratories Inc.		051565745	manufacture(37012-526)					

Revised: 2/2013

Shopko Stores Operating Co., LLC