LORATADINE AND PSEUDOEPHEDRINE SULFATE- loratadine and pseudoephedrine sulfate tablet, film coated, extended release Shopko Stores Operating Co., LLC

Drug Facts

ACTIVE INGREDIENTS (IN EACH TABLET)

Loratadine, USP 10 mg

Pseudoephedrine sulfate, USP 240 mg

PURPOSE

Antihistamine

Nasal decongestant

USES

- temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:
 - sneezing
 - itchy, watery eyes
 - runny nose
 - itching of the nose or throat
- reduces swelling of nasal passages
- temporarily relieves sinus congestion and pressure
- temporarily relieves nasal congestion due to the common cold, hay fever or other upper respiratory allergies
- temporarily restores freer breathing through the nose

WARNINGS

Do not use

- if you have ever had an allergic reaction to this product or any of its ingredients
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

Ask a doctor before use if you have

- heart disease
- thyroid disease
- high blood pressure
- diabetes
- trouble urinating due to an enlarged prostate gland
- 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.
- symptoms do not improve within 7 days or are accompanied by a fever
- nervousness, dizziness or sleeplessness occurs

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

• do not divide, crush, chew or dissolve the tablet

adults and children 12 years and	1 tablet daily with a full glass of water; not more than 1 tablet in
over	24 hours
children under 12 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

OTHER INFORMATION

- **sodium:** contains 10 mg/tablet
- **calcium:** contains 25 mg/tablet
- TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.
- store between 20° C to 25° C (68° F to 77° F).
- protect from light and store in a dry place

INACTIVE INGREDIENTS

Calcium carbonate, colloidal silicon dioxide, hydroxypropyl cellulose, hypromellose, iron oxide black, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch, propylene glycol, shellac glaze, sodium alginate, sodium citrate, talc and titanium dioxide

QUESTIONS?

Call 1-800-406-7984

PRINCIPAL DISPLAY PANEL SHOPKO[®] [†]Compare to the active ingredients of Claritin-D[®]24 Hour Original Prescription Strength 24 Hour Allergy Relief-D Loratadine, USP 10 mg/Antihis tamine Pseudoephedrine Sulfate, USP 240 mg/Nasal Decongestant Non-Drowsy*

Indoor & Outdoor Allergies

Relief of:

- Nasal and Sinus Congestion
- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Throat or Nose Due to Allergies

^{*}When taken as directed. See Drug Facts Panel.

Manufactured by: Ohm Laboratories Inc. 5093688/R0512



LORATADINE AND PSEUDOEPHEDRINE SULFATE loratadine and pseudoephedrine sulfate tablet, film coated, extended release **Product Information** HUMAN OTC DRUG NDC:37012-724 **Product Type** Item Code (Source) ORAL **Route of Administration Active Ingredient/Active Moiety Ingredient** Name **Basis of Strength** Strength LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN) LORATADINE 10 mg PSEUDO EPHEDRINE SULFATE (UNII: Y9DL7QPE6B) (PSEUDO EPHEDRINE -PSEUDOEPHEDRINE 240 mg UNII:7CUC9DDI9F) SULFATE **Inactive Ingredients Ingredient Name** Strength

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CALCIUM CARBONATE (UNII: H0 G9 379 FGK)	
COLLOIDAL SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HYDROXYPROPYL CELLULOSE (UNII: RFW2ET671P)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FERROSOFERRIC OXIDE (UNII: XM0 M87F357)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
PO VIDO NE (UNII: FZ989GH94E)	
STARCH, PREGELATINIZED CORN (UNII: 08232NY3SJ)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SHELLAC (UNII: 46N107B710)	
SODIUM ALGINATE (UNII: C269C4G2ZQ)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)	

Product Characteristics				
Color	white (White to Off-White)	Score	no score	
Shape	CAPSULE	Size	17mm	
Flavor		Imprint Code	RX724	
Contains				

Packaging

# Item Code	Package Description	Marketing Start Date	Marketing End Date				
1 NDC:37012-724-69	10 in 1 BLISTER PACK						
Marketing Information							
Marketing Info	rmation						
Marketing Info Marketing Category	rmation Application Number or Monogra	aph Citation Marketing Stat	rt Date Marketing End Date				
•		aph Citation Marketing Star 11/17/2004	rt Date Marketing End Date				

Labeler - Shopko Stores Operating Co., LLC (023252638)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	manufacture(37012-724)

Revised: 2/2013

Shopko Stores Operating Co., LLC