LORATADINE AND PSEUDOEPHEDRINE SULFATE- loratadine and pseudoephedrine sulfate tablet, film coated, extended release
HEB

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 over	1 tablet daily with a full glass of water; not more than 1 tablet in 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

Compare to Claritin-D®24 Hour active ingredients**

NDC 37808-0724-69

H-E-B®

ALLERGY RELIEF-D

Non-Drowsy*- 24 Hour Formulation

Loratadine, USP 10 mg/Antihistamine

Pseudoephedrine Sulfate, USP 240 mg/Nasal Decongestant

Indoor & Outdoor Allergies

Allergy & Congestion

Original Prescription Strength

Relief of:

- **Nasal & Sinus Congestion Due to Colds or Allergies**
- **Sneezing; Runny Nose; Itchy, Watery Eyes;**

Itchy Throat or Nose Due to Allergies

10 Extended-Release Tablets

***When taken as directed. See Drug Facts Panel.**

DISTRIBUTED BY H-E-B

5088127/R0711


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H-E-B
ALLERGY RELIEF–D
Non-Drowsy* - 24 Hour Formulation
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*When taken as directed. See Drug Facts Panel.
actual size
10
Extended-Release Tablets
5088127
5088127
Drug Facts
Active ingredients (in each tablet)
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Antihistamine
Pseudoephedrine sulfate, USP 240 mg
Nasal decongestant
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■ sneezing ■ itchy, watery eyes ■ runny nose ■ itching of the nose or throat
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Drug Facts (continued)
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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
Keep the carton. It contains important information. See end panel for expiration date.
Batch No.
Expiration Date:
Non Varnish Area
**The product is not manufactured or distributed by Schering-Plough Healthcare Products, Inc. CLARITIN-D® 24 HOUR is a registered trademark of Schering Corporation. R0711
MADE WITH PRIDE & CARE FOR H-E-B, SAN ANTONIO, TX 78204
GUARANTEE
We believe the high quality of this H-E-B product makes it an outstanding value. We hope you'll agree. If not, we'll cheerfully refund your money. Thanks for shopping with us.
0 41220 98494 7

LORATADINE AND PSEUDOEPHEDRINE SULFATE

loratadine and pseudoephedrine sulfate tablet, film coated, extended release

Product Information			
Product Type	HUMAN OTC DRUG LABEL	**Item Code (Source)**	NDC:37808-724
Route of Administration	ORAL	**DEA Schedule**	

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LORATADINE (LORATADINE)	LORATADINE	10 mg
PSEUDOEPHEDRINE SULFATE (PSEUDOEPHEDRINE)	PSEUDOEPHEDRINE SULFATE	240 mg

Inactive Ingredients	
Ingredient Name	**Strength**
CALCIUM CARBONATE	
COLLOIDAL SILICON DIOXIDE	
HYDROXYPROPYL CELLULOSE	
HYPROMELLOSES	
FERROSOFERRIC OXIDE	
LACTOSE MONOHYDRATE	
MAGNESIUM STEARATE	
CELLULOSE, MICROCRYSTALLINE	
POLYETHYLENE GLYCOLS	
POVIDONE	
STARCH, PREGELATINIZED CORN	
PROPYLENE GLYCOL	
SHELLAC	
SODIUM ALGINATE	
SODIUM CITRATE	
TALC	
TITANIUM DIOXIDE	

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:37808-724-69	10 in 1 BLISTER PACK		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076557	11/17/2004	

Labeler - HEB (007924756)

Registrant - Ranbaxy Pharmaceuticals Inc. (937890044)

Establishment

Name	Address	ID/FEI	Business Operations
Ohm Laboratories Inc.		051565745	manufacture(37808-724)

Revised: 9/2012 HEB