

LORATADINE- loratadine capsule, liquid filled
KROGER COMPANY

KROGER 686T

LORATADINE 10MG

DO NOT USE IF YOU HAVE EVER HAD AN ALLERGIC REACTION TO THIS PRODUCT OR ANY OF ITS INGREDIENTS.

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.

ANTI-HISTAMINE

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

IN CASE OF OVERDOSE, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

ADULTS AND CHILDREN 6 YEARS AND OVER: 1 CAPSULE DAILY; NOT MORE THAN 1 CAPSULE IN 24 HOURS.

CHILDREN UNDER 6 YEARS OF AGE: ASK A DOCTOR.

CONSUMERS WITH LIVER OR KIDNEY DISEASE: ASK A DOCTOR.

STORE BETWEEN 20-25 DEGREES CELSIUS (67-77 DEGREES FAHRENHEIT)

PROTECT FROM FREEZING

FD&C BLUE #1, GELATIN, MONO AND DIGLYCERIDE OF CAPRYLIC/CAPRIC ACID, PHARMACEUTICAL INK, POLYSORBATE 80, POVIDONE, PURIFIED WATER, SORBITOL SORBITAN SOLUTION.

LORATADINE

When Taken as Directed
Loratadine Capsules, 10 mg
24 HOUR

Kroger
NON-DROWSY*
Allergy Relief

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Allergy Relief
Loratadine Capsules, 10 mg
Antihistamine

COMPARE TO the active ingredient of CLARITIN[®] LIQUID-GELS[®]

NON-DROWSY*
Allergy Relief
Loratadine Capsules, 10 mg
Antihistamine

Relieves:

- Sneezing
- Itchy, Watery Eyes
- Itchy Throat or Nose
- Runny Nose

Indoor & Outdoor Allergies

24 HOUR

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SAFETY SEALED: DO NOT USE IF THE INDIVIDUAL BLISTER UNIT IMPRINTED WITH "LORATADINE LIQUID FILLED CAPSULES" IS OPEN OR TORN

READ AND KEEP CARTON FOR COMPLETE WARNINGS AND INFORMATION

Lot #:Exp. Date:

086749174-12-2013 35178-2

DISTRIBUTED BY THE KROGER CO.
CINCINNATI, OH 45202
MADE IN INDIA

QUALITY GUARANTEE
www.kroger.com

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Drug Facts	Purpose
Active Ingredient (in each capsule) ... Loratadine	Antihistamine
Uses	• To help relieve the symptoms of: <ul style="list-style-type: none">• hay fever• other allergic respiratory allergies• temporary relief of itching of the nose or throat
Warnings	• Do not use if you have ever had an allergic reaction to loratadine. • Ask a doctor before use if you have liver or kidney disease. Your doctor should determine if you should use this product. • When using this product, do not take more than that directed. Taking more than directed may cause drowsiness, dizziness, or other side effects. • Stop use and ask a doctor if an allergic reaction to this product occurs. Seek medical help right away if you experience a severe allergic reaction. • If pregnant or breast-feeding, ask a health professional before use.
Other Information	• Store between 20-25°C (68-77°F) • Protect from moisture
Inactive Ingredients (Pillar #1)	• croscarmellose sodium • polyethylene glycol 400 • hydroxypropyl cellulose • polydioxanone • polyethylene glycol 6000 • polyethylene glycol 400 • polyethylene glycol 6000 • polyethylene glycol 400 • polyethylene glycol 6000
Questions or comments?	• Visit us at: www.kroger.com

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41226-686
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LORATADINE (UNII: 7AJ03BO7QN) (LORATADINE - UNII:7AJ03BO7QN)	LORATADINE	10 mg

Inactive Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
GELATIN (UNII: 2G86QN327L)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE (UNII: FZ989GH94E)	
WATER (UNII: 059QF0K00R)	
SORBITOL (UNII: 506T60A25R)	
CAPRYLIC/CAPRIC MONO/DIGLYCERIDES (UNII: U72Q2I8C85)	

Product Characteristics

Color	blue	Score	no score
Shape	OVAL	Size	3mm
Flavor		Imprint Code	21
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41226-686-30	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	04/26/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA206214	04/26/2017	

Labeler - KROGER COMPANY (006999528)

Registrant - TIME CAP LABORATORIES INC (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
MARKSANS PHARMA LIMITED		925822975	manufacture(41226-686)

Revised: 5/2018

KROGER COMPANY