LORATADINE- loratadine capsule, liquid filled
SMART SENSE (KMART)

KMART 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.

ANTIHISTAMINE

TEMPORARILY RELIEVES THESE SYMPTOMS DUE TO HAY FEVER OR OTHR 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

loratadine capsule, liquid filled

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49738-686
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LORATADINE (UNII: 7AJO3BO7QN) (LORATADINE - UNII:7AJO3BO7QN)	LORATADINE	10 mg

Inactive Ingredients	
Ingredient Name	**Strength**
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GELATIN (UNII: 2G86QN327L)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
POVIDONE (UNII: FZ989GH94E)	
WATER (UNII: 059QF0KO0R)	
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:49738-686-07	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	01/31/2017	
2	NDC:49738-686-30	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	01/31/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA206214	01/31/2017	

Labeler - SMART SENSE (KMART) (008965873)

Registrant - TIME CAP LABORATORIES INC (037052099)

Establishment

Name	Address	ID/FEI	Business Operations
MARKSANS PHARMA LIMITED		925822975	manufacture(49738-686)