SUNMARK LORATADINE- loratadine tablet Strategic Sourcing Services LLC

McKesson Loratadine Tablets 10 mg Drug Facts

Active ingredient (in each tablet)

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

Other information

- do not use if printed foil under cap is broken or missing
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

lactose monohydrate, magnesium stearate, povidone, pregelatinized starch

Questions or comments?

1-800-719-9260

Principal Display Panel

COMPARE TO CLARITIN® TABLETS ACTIVE INGREDIENT

24 HOUR

loratadine tablets, 10 mg

Antihistamine

24 hour relief of sneezing; runny nose;

itchy, watery eyes;

itchy throat or nose

Indoor & Outdoor Allergies

ORIGINAL PRESCRIPTION STRENGTH

NON-DROWSY*

Actual Size

*When taken as directed.

See Drug Facts Panel.

GLUTEN FREE

70 TABLETS



SUNMARK LORATADINE

loratadine tablet

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:49348-818

Route of Administration ORAL

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

Inactive Ingredients			
Ingredient Name	Strength		
LACTO SE MO NO HYDRATE (UNII: EWQ57Q8 I5X)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
PO VIDONE, UNSPECIFIED (UNII: FZ989GH94E)			

Product Characteristics						
Color	WHITE	Score	no score			
Shape	OVAL	Size	8 mm			
Flavor		Imprint Code	L612			
Contains						

Packaging							
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:49348-818-01	10 in 1 CARTON	06/25/2007				
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product					
2	NDC:49348-818-44	30 in 1 CARTON	06/21/2007				
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product					
3	NDC:49348-818-12	1 in 1 CARTON	07/13/2007	06/07/2015			
3		60 in 1 BOTTLE; Type 0: Not a Combination Product					
4	NDC:49348-818-13	1 in 1 CARTON	06/21/2007				
4		90 in 1 BOTTLE; Type 0: Not a Combination Product					
5	NDC:49348-818-56	1 in 1 CARTON	03/31/2014				
5		70 in 1 BOTTLE; Type 0: Not a Combination Product					
6	NDC:49348-818-45	1 in 1 CARTON	10/24/2018				
6		30 in 1 BOTTLE; Type 0: Not a Combination Product					

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
ANDA	ANDA076301	06/21/2007			

Labeler - Strategic Sourcing Services LLC (116956644)

Revised: 3/2020 Strategic Sourcing Services LLC