

LORATADINE ALLERGY RELIEF- loratadine tablet
Asclemed USA, Inc.

Loratadine Allergy Relief

Drug Facts

Active ingredient (in each tablet)

Loratadine, USP 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	ask a doctor

age

ask a doctor

consumers with liver or
kidney disease

ask a doctor

Other Information

- store between 20° to 25° C (68° to 77° F)
- protect from excessive moisture
- **TAMPER EVIDENT: DO NOT USE IF BLISTER UNITS ARE TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING.**

Inactive ingredients

corn starch, lactose monohydrate, magnesium stearate, pregelatinized starch

Questions?

call **1-800-406-7984**

Relabeled and Repackaged by:

Enovachem PHARMACEUTICALS

Torrance, CA 90501

PRINCIPAL DISPLAY PANEL

Relabeled By:
 370 Van Ness Ave.
 Suite 1425-1408
 Torrance, CA 90501
Loratadine Tablets, USP 10 mg
 NDC: 76420-330-11
 Qty: 10
 Distributed By: Ohm Laboratories Inc.
 Source NDC: 51660-526-11
 Description: white to off white, round/RX526 In blister pack
 Lot #: 00000000
 Exp: (1) 03/76420 33011 0
 (17)
 (10) 00000000
 (21)
 Batch #: 00000000
 Drug Status: OTC
 CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT.
 KEEP OUT OF REACH OF CHILDREN. STORE AT 20-25C (68-77F) (SEE USP CONTROLLED ROOM TEMP).

Loratadine Tablets, USP 10 mg
 NDC: 76420-330-11
 S/N:
 Qty: 10
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-11
 S/N:
 Qty: 10
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-11
 S/N:229939
 Qty: 10

Relabeled By:
 370 Van Ness Ave.
 Suite 1425-1408
 Torrance, CA 90501
Loratadine Tablets, USP 10 mg
 NDC: 76420-330-05
 Qty: 500
 Distributed By: Ohm Laboratories Inc.
 Source NDC: 51660-526-05
 Description: white to off white, round/RX526
 Lot #: 00000000
 Exp: (1) 03/76420 33005 9
 (17)
 (10) 00000000
 (21)
 Batch #: 00000000
 Drug Status: OTC
 CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT.
 KEEP OUT OF REACH OF CHILDREN. STORE AT 20-25C (68-77F) (SEE USP CONTROLLED ROOM TEMP).

Loratadine Tablets, USP 10 mg
 NDC: 76420-330-05
 S/N:
 Qty: 500
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-05
 S/N:
 Qty: 500
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-05
 S/N:
 Qty: 500

Relabeled By:
 370 Van Ness Ave.
 Suite 1425-1408
 Torrance, CA 90501
Loratadine Tablets, USP 10 mg
 NDC: 76420-330-30
 Qty: 30
 Distributed By: Ohm Laboratories Inc.
 Source NDC: 51660-526-30
 Description: white to off white, round/RX526
 Lot #: 00000000
 Exp: (1) 03/76420 33030 1
 (17)
 (10) 00000000
 (21)
 Batch #: 00000000
 Drug Status: OTC
 CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT.
 KEEP OUT OF REACH OF CHILDREN. STORE AT 20-25C (68-77F) (SEE USP CONTROLLED ROOM TEMP).

Loratadine Tablets, USP 10 mg
 NDC: 76420-330-30
 S/N:
 Qty: 30
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-30
 S/N:
 Qty: 30
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-30
 S/N:
 Qty: 30

Relabeled By:
 370 Van Ness Ave.
 Suite 1425-1408
 Torrance, CA 90501
Loratadine Tablets, USP 10 mg
 NDC: 76420-330-10
 Qty: 10
 Distributed By: Ohm Laboratories Inc.
 Source NDC: 51660-526-XX
 Description: white to off white, round/RX526
 Lot #: 00000000
 Exp: (1) 03/76420 33010 3
 (17)
 (10) 00000000
 (21)
 Batch #: 00000000
 Drug Status: OTC
 CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT.
 KEEP OUT OF REACH OF CHILDREN. STORE AT 20-25C (68-77F) (SEE USP CONTROLLED ROOM TEMP).

Loratadine Tablets, USP 10 mg
 NDC: 76420-330-10
 S/N:
 Qty: 10
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-10
 S/N:
 Qty: 10
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-10
 S/N:
 Qty: 10

Relabeled By:
 370 Van Ness Ave.
 Suite 1425-1408
 Torrance, CA 90501
Loratadine Tablets, USP 10 mg
 NDC: 76420-330-31
 Qty: 30
 Distributed By: Ohm Laboratories Inc.
 Source NDC: 51660-526-31
 Description: white to off white, round/RX526 In blister pack
 Lot #: 00000000
 Exp: (1) 03/76420 33031 8
 (17)
 (10) 00000000
 (21)
 Batch #: 00000000
 Drug Status: OTC
 CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT.
 KEEP OUT OF REACH OF CHILDREN. STORE AT 20-25C (68-77F) (SEE USP CONTROLLED ROOM TEMP).

Loratadine Tablets, USP 10 mg
 NDC: 76420-330-31
 S/N:
 Qty: 30
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-31
 S/N:
 Qty: 30
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-31
 S/N:
 Qty: 30

Relabeled By:
 370 Van Ness Ave.
 Suite 1425-1408
 Torrance, CA 90501
Loratadine Tablets, USP 10 mg
 NDC: 76420-330-60
 Qty: 60
 Distributed By: Ohm Laboratories Inc.
 Source NDC: 51660-526-XX
 Description: white to off white, round/RX526
 Lot #: 00000000
 Exp: (1) 03/76420 33060 8
 (17)
 (10) 00000000
 (21)
 Batch #: 00000000
 Drug Status: OTC
 CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT.
 KEEP OUT OF REACH OF CHILDREN. STORE AT 20-25C (68-77F) (SEE USP CONTROLLED ROOM TEMP).

Loratadine Tablets, USP 10 mg
 NDC: 76420-330-60
 S/N:
 Qty: 60
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-60
 S/N:
 Qty: 60
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-60
 S/N:
 Qty: 60

Relabeled By:
 370 Van Ness Ave.
 Suite 1425-1408
 Torrance, CA 90501
Loratadine Tablets, USP 10 mg
 NDC: 76420-330-01
 Qty: 100
 Distributed By: Ohm Laboratories Inc.
 Source NDC: 51660-526-01
 Description: white to off white, round/RX526
 Lot #: 00000000
 Exp: (1) 03/76420 33001 1
 (17)
 (10) 00000000
 (21)
 Batch #: 00000000
 Drug Status: OTC
 CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT.
 KEEP OUT OF REACH OF CHILDREN. STORE AT 20-25C (68-77F) (SEE USP CONTROLLED ROOM TEMP).

Loratadine Tablets, USP 10 mg
 NDC: 76420-330-01
 S/N:
 Qty: 100
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-01
 S/N:
 Qty: 100
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-01
 S/N:
 Qty: 100

Relabeled By:
 370 Van Ness Ave.
 Suite 1425-1408
 Torrance, CA 90501
Loratadine Tablets, USP 10 mg
 NDC: 76420-330-90
 Qty: 90
 Distributed By: Ohm Laboratories Inc.
 Source NDC: 51660-526-XX
 Description: white to off white, round/RX526
 Lot #: 00000000
 Exp: (1) 03/76420 33090 5
 (17)
 (10) 00000000
 (21)
 Batch #: 00000000
 Drug Status: OTC
 CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT.
 KEEP OUT OF REACH OF CHILDREN. STORE AT 20-25C (68-77F) (SEE USP CONTROLLED ROOM TEMP).

Loratadine Tablets, USP 10 mg
 NDC: 76420-330-90
 S/N:
 Qty: 90
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-90
 S/N:
 Qty: 90
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-90
 S/N:
 Qty: 90

Relabeled By:
 370 Van Ness Ave.
 Suite 1425-1408
 Torrance, CA 90501
Loratadine Tablets, USP 10 mg
 NDC: 76420-330-03
 Qty: 300
 Distributed By: Ohm Laboratories Inc.
 Source NDC: 51660-526-53
 Description: white to off white, round/RX526
 Lot #: 00000000
 Exp: (1) 03/76420 33003 5
 (17)
 (10) 00000000
 (21)
 Batch #: 00000000
 Drug Status: OTC
 CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT.
 KEEP OUT OF REACH OF CHILDREN. STORE AT 20-25C (68-77F) (SEE USP CONTROLLED ROOM TEMP).

Loratadine Tablets, USP 10 mg
 NDC: 76420-330-03
 S/N:
 Qty: 300
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-03
 S/N:
 Qty: 300
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-03
 S/N:
 Qty: 300

Relabeled By:
 370 Van Ness Ave.
 Suite 1425-1408
 Torrance, CA 90501
Loratadine Tablets, USP 10 mg
 NDC: 76420-330-20
 Qty: 20
 Distributed By: Ohm Laboratories Inc.
 Source NDC: 51660-526-XX
 Description: white to off white, round/RX526
 Lot #: 00000000
 Exp: (1) 03/76420 33020 2
 (17)
 (10) 00000000
 (21)
 Batch #: 00000000
 Drug Status: OTC
 CAUTION: FEDERAL LAW PROHIBITS DISPENSING WITHOUT PRESCRIPTION. SEE PACKAGE INSERT.
 KEEP OUT OF REACH OF CHILDREN. STORE AT 20-25C (68-77F) (SEE USP CONTROLLED ROOM TEMP).

Loratadine Tablets, USP 10 mg
 NDC: 76420-330-20
 S/N:
 Qty: 20
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-20
 S/N:
 Qty: 20
 Loratadine Tablets, USP 10 mg
 NDC: 76420-330-20
 S/N:
 Qty: 20

LORATADINE ALLERGY RELIEF

loratadine tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76420-330(NDC:51660-526)
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
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	

MAGNESIUM STEARATE (UNII: 70097M6I30)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	white (White to Off White)	Score	no score
Shape	ROUND	Size	6mm
Flavor		Imprint Code	RX526
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76420-330-10	10 in 1 BOTTLE; Type 0: Not a Combination Product	04/19/2025	
2	NDC:76420-330-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	04/19/2025	
3	NDC:76420-330-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	04/19/2025	
4	NDC:76420-330-90	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/19/2025	
5	NDC:76420-330-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	04/19/2025	
6	NDC:76420-330-03	300 in 1 BOTTLE; Type 0: Not a Combination Product	04/19/2025	
7	NDC:76420-330-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	04/19/2025	
8	NDC:76420-330-31	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	04/19/2025	
9	NDC:76420-330-11	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	04/19/2025	
10	NDC:76420-330-20	20 in 1 BOTTLE; Type 0: Not a Combination Product	05/21/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076134	11/01/2017	

Labeler - Asclemed USA, Inc. (059888437)

Establishment

Name	Address	ID/FEI	Business Operations
ASCLEMED USA INC. DBA ENOVACHEM		059888437	relabel(76420-330) , repack(76420-330)