LEVOCETIRIZINE DIHYDROCHLORIDE- levocetirizine dihydrochloride tablet, coated

Rite Aid Corporation

Levocetirzine Dihydrochloride Tablets USP, 5 mg

Active ingredient (in each tablet)

Levocetirizine dihydrochloride USP, 5 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other respiratory allergies:

- runny nose
- sneezing
- itchy, watery eyes
- itching of the nose or throat

Warnings

Do not use

- if you have kidney disease
- if you have ever had an allergic reaction to this product or any of its ingredients or to an antihistamine containing cetirizine

Ask a doctor before use if you have

· ever had trouble urinating or emptying your bladder

When using this product

- drowsiness may occur
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery

Stop use and ask doctor if

- you have trouble urinating or emptying your bladder
- an allergic reaction to this product occurs. Seek medical help right away.

If pregnant or breast-feeding:

• if breast-feding: not recommended

• if pregnant: 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 65 years of age and older	ask a doctor
adults and children 12-64 years of age	 take 1 tablet (5 mg) once daily in the evening do not take more than 1 tablet (5 mg) in 24 hours 1/2 tablet (2.5 mg) once daily in the evening may be appropriate for less severe symptoms
children 6-11 years of age	 take 1/2 tablet (2.5 mg) once daily in the evening do not take more than 1/2 tablet (2.5 mg) in 24 hours
children under 6 years of age	• do not use
consumers with kidney disease	• do not use

Other information

- store between 20° and 25°C (68° and 77°F)
- (bottles only) safety sealed: do not use if carton was opened or if printed foil inner seal on bottle is torn or missing
- (blister only) safety sealed: do not use if seal is broken or if individual blister unit is open or torn

Inactive ingredients

colloidal silicon dioxide, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide

Questions or comments?

Call **1-888-375-3784**

PRINCIPAL DISPLAY PANEL



ORIGINAL **PRESCRIPTION** STRENGTH

allergy relief

levocetirizine dihydrochloride tablets, 5 mg

antihistamine

120 TABLETS 5 ma EACH Active ingredient (in each tablet) Purpose Levocetirizine dihydrochloride USP, 5 mg......Antihistamine Uses temporarily relieves these symptoms due to hav fever or other respiratory allergies: ■ runny nose ■ sneezing ■ itchy, watery eyes ■ itching of the nose or throat Warnings Do not use ■ if you have kidney disease if you have ever had an allergic reaction to this product or any of its ingredients or to an antihistamine containing cetirizine Ask a doctor before use if you have ■ ever had trouble urinating or emptying your bladder When using this product ■ drowsiness may occur ■ avoid alcoholic drinks ■ alcohol, sedatives, and tranquilizers may increase drowsiness ■ be careful when driving a motor vehicle or operating machinery Stop use and ask a doctor if \(\bigsi \) you have trouble urinating or emptying your bladder ■ an allergic reaction to this product occurs. Seek medical help right away. If pregnant or breast-feeding: ■ if breast-feeding: not recommended if pregnant; ask a health professional before use

Peel Here

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DISTRIBUTED BY: RITE AID 30 HUNTER LANE CAMP HILL, PA 17011 MADE IN INDIA

EXP

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LEVOCETIRIZINE DIHYDROCHLORIDE

ORAL

levocetirizine dihydrochloride tablet, coated

Product Information

Route of Administration

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11822-5252(NDC:43598-735)

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength		
	levocetirizine dihydrochloride	5 mg		

Inactive Ingredients				
Ingredient Name	Strength			
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)				
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				

Polyethylene Glycol, Unspecified (UNII: 3WO0SDWIA)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	

Product Characteristics			
Color	white	Score	2 pieces
Shape	OVAL	Size	9mm
Flavor		Imprint Code	L
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:11822- 5252-7	1 in 1 CARTON	02/15/2020	
1		35 in 1 BOTTLE; Type 0: Not a Combination Product		
	NDC:11822- 5252-1	1 in 1 CARTON	02/15/2020	
2		120 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	ANDA210375	02/15/2020	

Labeler - Rite Aid Corporation (014578892)

Establishment			
Name	Address	ID/FEI	Business Operations
Dr.Reddy's Laboratories Limited-FTO 3		918608162	analysis(11822-5252), manufacture(11822-5252)

Establishment				
Name	Address	ID/FEI	Business Operations	
Reed-Lane, Inc.		001819879	repack(11822-5252)	

Revised: 11/2019 Rite Aid Corporation