

LEVOCETIRIZINE DIHYDROCHLORIDE- levocetirizine dihydrochloride tablet, coated
Wal-Mart Stores Inc

Drug Facts

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-feeding: 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	<ul style="list-style-type: none">• ask a doctor
adults and children 12-64 years of age	<ul style="list-style-type: none">• 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	<ul style="list-style-type: none">• 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	<ul style="list-style-type: none">• do not use
consumers with kidney disease	<ul style="list-style-type: none">• do not use

Other information

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

Inactive ingredients

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

Questions or comments?

Call 1-888-375-3784

Carton Label

No Coating Area



equate™

Allergy Relief

Levocetirizine Dihydrochloride Tablets USP, 5 mg

Antihistamine

ORIGINAL PRESCRIPTION STRENGTH

Drug Facts (continued)

Other information

- store between 20° and 25°C (68° and 77°F)
- safety sealed; do not use if carton was opened or if printed foil inner seal on bottle is torn or missing

Inactive ingredients colloidal silicon dioxide, hydroxypropyl methylcellulose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide

Questions or comments? 1-888-287-1915

Read directions and warnings before use. Keep this carton. It has important information.



Satisfaction guaranteed - Or we'll replace it or give you your money back. For questions or comments or to report an undesired reaction or side effect, please call 1-888-287-1915.

DISTRIBUTED BY: Walmart Inc., Bentonville, AR 72716
PRODUCT OF INDIA
 *This product is not manufactured or distributed by Chatterm, Inc. (part of the Sanofi Group), distributor of Xyzal® Allergy 24HR Tablets. Xyzal® is a registered trademark of UCB Group of companies.



equate™

- Clinically proven relief in 60 minutes
- Clinically proven relief from indoor and outdoor allergies
- Clinically proven relief for 24 hours



Actual Size

Revised: 06/19

27/32

equate™

NDC 4905-992-35



Allergy Relief

Levocetirizine Dihydrochloride Tablets USP, 5 mg

Antihistamine

ORIGINAL PRESCRIPTION STRENGTH

Relief of:

- Sneezing
- Runny nose
- Itchy nose or throat
- Itchy, watery eyes



Actual Size

5 mg 35 TABLETS

Drug Facts

Active ingredient (in each tablet) Purpose
Levocetirizine dihydrochloride USP, 5 mg.....Antihistamine

Uses temporarily relieves these symptoms due to hay fever or other respiratory allergies:
 ■ runny nose ■ itchy, watery eyes
 ■ sneezing ■ 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

■ 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

Keep out of reach of children.

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Directions

adults 65 years of age and older	■ ask a doctor
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children under 6 years of age	■ do not use
consumers with kidney disease	■ do not use



No Coating Area



Blister Carton



LEVOCETIRIZINE DIHYDROCHLORIDE

levocetirizine dihydrochloride tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-492(NDC:43598-735)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
levocetirizine dihydrochloride (UNII: SOD6A38AGA) (levocetirizine - UNII:6U5EA9RT2O)	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)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	

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
1	NDC:49035-492-35	1 in 1 BOTTLE	07/26/2019	
1		35 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:49035-492-79	2 in 1 BLISTER PACK	07/26/2019	
2		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210375	07/26/2019	

Labeler - Wal-Mart Stores Inc (051957769)

Revised: 7/2019

Wal-Mart Stores Inc