

ALLERGY RELIEF- levocetirizine dihydrochloride tablet, coated
AMERISOURCEBERGEN DRUG CORPORATION

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 eveningdo not take more than 1 tablet (5 mg) in 24 hours1/2 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 eveningdo 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

Compare to Xyzal® Allergy 24HR
active ingredient*

Good
Neighbor

Pharmacy®

ORIGINAL PRESCRIPTION STRENGTH

Allergy Relief
levocetirizine dihydrochloride
tablets USP, 5 mg
Antihistamine

Relief of:

- Sneezing
- Runny Nose
- Itchy Nose or Throat
- Itchy, Watery Eyes

24 HOUR



2 1 7 5 6 0 0 5 1



NDC 46122-531-61

ORIGINAL PRESCRIPTION STRENGTH

Allergy Relief

levocetirizine dihydrochloride tablets USP, 5 mg

Antihistamine

Drug Facts	Drug Facts (continued)
Active ingredient (in each tablet) levocetirizine dihydrochloride USP, 5 mg...Antihistamine	Directions ask a doctor
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	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
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	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
Ask a doctor before use if you have: • ever had trouble urinating or emptying your bladder	children under 6 years of age • do not use
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	consumers with kidney disease • do not use
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.	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
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).	Inactive ingredients colloidal silicon dioxide, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, titanium dioxide
ISSUED: 08/23	Questions or comments? Call 1-888-376-3784

This product is not manufactured or distributed by Chatter, Inc. (part of the Sanofi Group), distributor of Xyzal® Allergy 24HR Tablets. Xyzal® is a registered trademark of UCB Group of companies.

Made in India

Distributed By
AmerisourceBergen
1 West First Avenue
Conshohocken, PA 19428
Questions or Concerns?
www.gnp.org



Compare to Xyzal® Allergy 24HR active ingredient*

NDC 46122-531-61

ORIGINAL PRESCRIPTION STRENGTH

Allergy Relief

levocetirizine dihydrochloride tablets USP, 5 mg

Antihistamine

Relief of:

- Sneezing
- Runny Nose
- Itchy Nose or Throat
- Itchy, Watery Eyes

24 HOUR

actual size

80 tablets



NDC 46122-531-61

ORIGINAL PRESCRIPTION STRENGTH

Allergy Relief

levocetirizine dihydrochloride tablets USP, 5 mg

Antihistamine

80 tablets 24 hr

Actual Size

ABCF 10251800



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Bottle Label


Good
Neighbor
Pharmacy®

ORIGINAL PRESCRIPTION STRENGTH

Allergy Relief
levocetirizine
dihydrochloride
tablets USP, 5 mg

Antihistamine

24 HOUR



NDC 46122-531-61

ORIGINAL PRESCRIPTION STRENGTH

Allergy Relief

levocetirizine dihydrochloride tablets USP, 5 mg

Antihistamine

80 tablets

24 HOUR

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 ■ 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 ■ you have trouble urinating or

(Continued On Back Of Label)

Distributed By: AmerisourceBergen
1 West First Avenue
Conshohocken PA 19428
Made in India

ISSUED: 08/23

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LOT
EXP
PEEL HERE

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).

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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

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

Questions and comments? call 1-888-375-3784

ALLERGY RELIEF

levocetirizine dihydrochloride tablet, coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:46122-531
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:46122-531-05	1 in 1 CARTON	04/20/2018	
1		35 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:46122-531-52	2 in 1 CARTON	07/06/2018	
2		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:46122-531-61	1 in 1 CARTON	09/01/2023	
3		80 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	04/20/2018	

Labeler - AMERISOURCEBERGEN DRUG CORPORATION (007914906)

Revised: 8/2023

AMERISOURCEBERGEN DRUG CORPORATION