DG HEALTH ALL DAY ALLERGY- cetirizine hydrochloride tablet, film coated
Dolgencorp Inc

Dolgencorp, LLC All Day Allergy Drug Facts

Active ingredient (in each tablet)

Cetirizine HCl 10 mg

Purpose

Antihistamine

Uses

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

- runny nose
- sneezing
- itchy, watery eyes
- 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 or to an antihistamine containing hydroxyzine.

Ask a doctor before use if you have

liver or kidney disease. Your doctor should determine if you need a different dose.

Ask a doctor or pharmacist before use if you are

taking tranquilizers or sedatives.

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

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 and children 6 years and over	one 10 mg tablet once daily; do not take more than one 10 mg tablet in 24 hours. A 5 mg product may be appropriate for less severe symptoms.
adults 65 years and over	ask a doctor
children under 6 years of age	ask a doctor
consumers with liver or kidney disease	ask a doctor

Other information

- store between 20 25°C (68-77°F)
- Do not use if blister unit is broken or torn

Inactive ingredients

corn starch, FD&C blue no. 1 aluminum lake, hypromellose, lactose monohydrate, magnesium stearate, polydextrose, polyethylene glycol, povidone, titanium dioxide, triacetin

Questions or comments?

1-888-309-9030

Principal Display Panel

DG™|health

Compare to the active ingredient of Zyrtec®

Original Prescription Strength

All Day Allergy

Cetirizine Hydrochloride Tablets, 10 mg

Antihistamine

24 HOUR

Indoor & Outdoor Allergies

24 Hour Relief of:

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

14 Tablets

10 mg

Actual Tablet Size

Original Prescription Strength

All Day Allergy

Cetirizine Hydrochloride Tablets, 10 mg

Antihistamine

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DG™ health

Compare to the active ingredient of Zyrtec®*

Original Prescription Strength

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Cetirizine Hydrochloride Tablets, 10 mg

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24 HOUR

Indoor & Outdoor Allergies

24 Hour Relief of:

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

10 mg

14 Tablets

Actual Tablet Size

4H266 VT C2

CODE AREA

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Drug Facts (continued)

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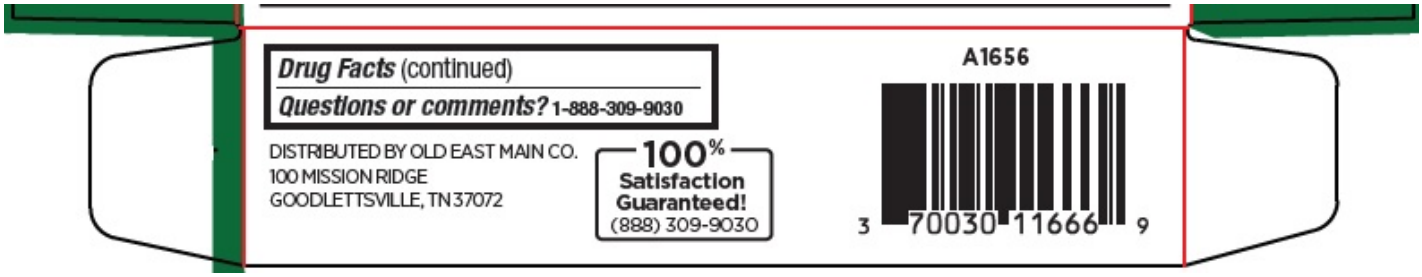
Tear along perforation, peel off paper and push product through foil. If difficult to open use scissors.

Indoor & Outdoor Allergy Relief

*This product is not manufactured or distributed by McNeil Consumer Healthcare, Division of McNeil-PPC Inc., distributor of Zyrtec®.



DG HEALTH ALL DAY ALLERGY

cetirizine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-458
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	10 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYDEXTROSE (UNII: VH2XOU12IE)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
TRIACETIN (UNII: XHX3C3X673)	

Product Characteristics

Color	WHITE	**Score**	no score
Shape	OVAL	**Size**	10mm
Flavor		**Imprint Code**	4H2
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-458-	1 in 1 PACKAGE	01/08/2010	

1	39	1 in 1 PACKAGE	01/08/2010	
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:55910-458-95	1 in 1 PACKAGE	10/10/2011	05/18/2015
2		45 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:55910-458-58	1 in 1 CARTON	05/08/2017	09/01/2021
3		40 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:55910-458-66	14 in 1 CARTON	06/16/2023	
4		1 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	ANDA078336	01/08/2010	

Labeler - Dolgencorp Inc (068331990)

Revised: 6/2023 Dolgencorp Inc