CETIRIZINE HCL- cetirizine hcl capsule
DOLGENCORP INC

Cetirizine HCl Capsules, 10 mg-dollar general

Active ingredient (in each capsule)

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
- alcohol, sedatives, and tranquilizers may increase drowsiness
- avoid alcoholic drinks
- 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.

Directions

adults and children 6 years and over	one 10 mg capsule once daily; do not take more than one 10 mg capsule 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 at 20º-25ºC (68º-77ºF)
- avoid high humidity and excessive heat above 40ºC (104ºF)
- protect from light

Inactive ingredients

FD&C yellow #6, gelatin, glycerin, mannitol, pharmaceutical ink, polyethylene glycol, purified water, sodium hydroxide, sorbitan, sorbitol

Questions or comments?

call toll free **1-888-235-2466** (Mon - Fri 9AM - 5PM EST)

†This product is not manufactured or distributed by the owners of Zyrtec ®

TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING

KEEP THIS CARD FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

DISTRIBUTED BY OLD EAST MAIN CO.

100 MISSION RIDGE

GOODLETTESVILLE, TN 37072

100% Satisfaction Gaurenteed

(888) 309-9030

Principal Display Panel-25's count

NDC 55910-784-86

DG Health

Compare to the

active ingredient

of Zyrtec

All-Day

Allergy Relief

Cetirizine HCl Capsules,

10mg/ Antihistamine

24 Hour Relief of:

Sneezing
Runny nose
Itchy, watery eyes
Itchy throat or nose

Indoor & Outdoor Allergies

25 Softgels**

**Liquid-filled capsules

Principal Display Panel- 40's count carton

NDC 55910-784-15

DG Health

Compare to the

active ingredient

of Zyrtec

All-Day

Allergy Relief

Cetirizine HCl Capsules,

10mg/ Antihistamine

24 Hour Relief of:

Sneezing
Runny nose
Itchy, watery eyes
Itchy throat or nose

Indoor & Outdoor Allergies

40 Softgels**

**Liquid-filled capsules

CETIRIZINE HCL

cetirizine hcl capsule

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-784
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
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GELATIN (UNII: 2G86QN327L)	
GLYCERIN (UNII: PDC6A3C0OX)	
MANNITOL (UNII: 3OWL53L36A)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SORBITAN (UNII: 6O92ICV9RU)	
SORBITOL (UNII: 506T60A25R)	

Product Characteristics

Color	orange	**Score**	no score
Shape	OVAL	**Size**	13mm
Flavor		**Imprint Code**	CE1
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-784-86	1 in 1 CARTON	11/25/2024	
1		25 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
2	NDC:55910-784-15	1 in 1 CARTON	11/25/2024	
2		40 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA022429	11/25/2024	

Labeler - DOLGENCORP INC (068331990)

Registrant - Bionpharma Inc. (079637826)

Establishment

Name	Address	ID/FEI	Business Operations
Patheon Softgels Inc.		002193829	manufacture(55910-784)

Revised: 11/2024 DOLGENCORP INC