ALL DAY ALLERGY RELIEF- cetirizine hydrochloride tablet DOLGENCORP, INC.

Dollar General Cetirizine Hydrochloride Tablets USP, 10 mg

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 useif 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 haveliver 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 ifan 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 adults 65 years and over children under 6 years of age consumers with liver or kidney

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.

ask a doctor

ask a doctor

ask a doctor

disease

Other information

store between 20° to 25°C (68° to 77°F).

Inactive ingredientscorn starch, hypromellose, lactose monohydrate, magnesium stearate, polyethylene glycol, povidone, titanium dioxide

Questions or comments? Call 1-888-309-9030





LOGG/LOD

RETAIN CARTON FOR COMPLETE PRODUCT INFORMATION

CODE NO.: GU/DRUGS/515

Drug Facts

Active ingredient **Purpose** (in each tablet)

Cetirizine HCI 10 mg. .Antihistamine

- 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

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 on 10 mg tablet in 24 hours. A 5 mg product may be appropriate for les severe symptoms.	
adults 65 years and over	ask a doctor	
children under 6 years of age	ask a doctor	
consumers with	ask a doctor	

Other information

liver or kidney disease

Drug Facts (continued)

Inactive ingredients

corn starch, hypromellose, lactose monohydrate, magnesium stearate, magnesium stearate, polyethylene glycol, povidone, titanium dioxide

Questions or comments? Call 1-888-309-9030

†This product is not manufactured or distributed by Johnson & Johnson Consumer Inc., McNeil Consumer Healtho Division, owner of the registered trademark Zyrtec® Tablets.

TAMPER EVIDENT: DO NOT USE THIS PRODUCT IF THE IMPRINTED FOIL SEAL OVER THE MOUTH OF THE BOTTLE IS CUT, TORN, BROKEN OR MISSING

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692R 1123

100%

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Compare to the active ingredient of Zyrtec* tablets t

45

Tablets

Varnish/Print Omit Area

ndoor & Outdoor Allergies

Varnish

ALL DAY ALLERGY RELIEF

cetirizine hydrochloride tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:55910-269

ORAL Route of Administration

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZ INE HYDROCHLORIDE	10 mg

Inactive Ingredients			
Ingredient Name	Strength		
HYPROMELLOSES (UNII: 3NXW29V3WO)			
MAGNESIUM STEARATE (UNII: 70097M6I30)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
POVIDONE (UNII: FZ 989GH94E)			
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
STARCH, CORN (UNII: O8232NY3SJ)			

Product Characteristics			
Color	white (White to off white)	Score	no score
Shape	RECTANGLE (Rounded-off, rectangular shaped tablet)	Size	9mm
Flavor		Imprint Code	J;220
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:55910-269- 58	1 in 1 CARTON	12/20/2023		
1		45 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	ANDA078933	12/20/2023	

Labeler - DOLGENCORP, INC. (068331990)

Registrant - TIME CAP LABORATORIES, INC. (037052099)

Establishment				
Name	Address	ID/FEI	Business Operations	
MARKSANS PHARMA LIMITED		925822975	manufacture(55910-269)	

Revised: 12/2023 DOLGENCORP, INC.