ALLERGY RELIEF- cetirizine hcl tablet A-S Medication Solutions

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
- 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 (1-800-222-1222) right away.

Directions

adults and children 6 years	Take 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
and over	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° to 25°C (68° to 77°F)
- contains no ingredient made from a gluten-containing grain(wheat, barley or rye)

Inactive ingredients

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

Questions or comments?

Call 1-888-588-1418 Monday-Friday 9AM-5PM EST

Principal Display Panel

*Compare to the active ingredient in $\mathbf{Zyrtec} \, \mathbb{R}$

Allergy Relief

Cetirizine HCl Tablets USP, 10 mg/Antihistamine

Indoor & Outdoor Allergies

24 hour relief of

- Sneezing
- Runny Nose
- Itchy, Watery Eyes

• Itchy Throat or Nose

Original Prescription Strength

Tablets

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

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

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

Distributed by:

Camber Consumer Care Inc, Piscataway, NJ 08854, USA

Cetirizine HCI



ALLERGY RELIEF

cetirizine hcl tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:50090-3906(NDC:69230-304)	
Route of Administration	ORAL			

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:Y07261ME24)	CETIRIZ INE HYDROCHLORIDE	10 mg

Inactive Ingredients				
Ingredient Name	Strength			
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)				
SILICON DIOXIDE (UNII: ETJ7Z 6XBU4)				
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)				
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				

Product Characteristics			
Color	white	Score	2 pieces
Shape	OVAL	Size	9mm
Flavor		Imprint Code	G;4
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50090- 3906-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	12/31/2018	
2	NDC:50090- 3906-2	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/31/2018	
3	NDC:50090- 3906-3	90 in 1 BOTTLE; Type 0: Not a Combination Product	12/31/2018	
4	NDC:50090- 3906-0	14 in 1 BOTTLE; Type 0: Not a Combination Product	12/31/2018	

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA209274	12/31/2018	

Labeler - A-S Medication Solutions (830016429)

Establishment				
Name	Address	ID/FEI	Business Operations	
A-S Medication Solutions		830016429	RELABEL(50090-3906), REPACK(50090-3906)	

Revised: 2/2023 A-S Medication Solutions