CETIRIZINE HYDROCHLORIDE- cetirizine hydrochloride tablet **SAFEWAY**

Indoor and Outdoor Allergies Allergy Relief

Cetrizine Hydrochloride Tablets, USP 10mg

Antihistamine

24 Hour Relief of: Sneezing Runny Nose Itchy, Watery Eyes Itchy Throat or Nose

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

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

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° 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

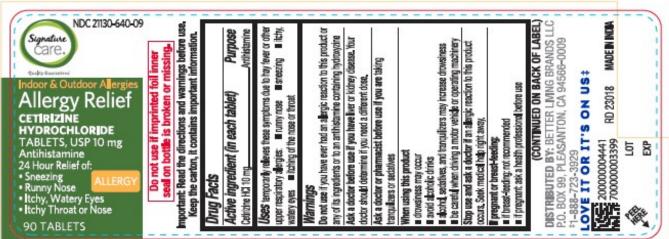
Questions?

call **1-877-770-3183** Mon-Fri 8:00 AM EST to 5:00 PM PST

Principal Display Panel



Principal Display Panel



Inside (adhesive side)



how2recycle.info

ASTIC **BOTTLE**

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Drug Facts (continued)



CETIRIZINE HYDROCHLORIDE

cetirizine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21130-640
Route of Administration	ORAL		

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	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)		

POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)

Product Characteristics			
Color	white (White to off-white)	Score	2 pieces
Shape	RECTANGLE (rounded off rectangular)	Size	9mm
Flavor		Imprint Code	G;4
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21130-640- 03	30 in 1 BOTTLE; Type 0: Not a Combination Product	06/30/2023	
2	NDC:21130-640- 18	180 in 1 CARTON; Type 0: Not a Combination Product	06/30/2023	
3	NDC:21130-640- 09	90 in 1 BOTTLE; Type 0: Not a Combination Product	06/30/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA209274	06/30/2023	

Labeler - SAFEWAY (009137209)

Revised: 12/2023 SAFEWAY