ALLERGY RELIEF- cetirizine hydrochloride tablet, film coated H E B

HEB Allergy Relief 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	one 10 mg tablet once daily; do not take more than one 10 mg tablet in 24
over	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	ask a doctor
disease	

Other information

- store between 20-25°C (68-77°F)
- do not use if printed foil under cap is broken or missing

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?

1-800-719-9260

Principal Display Panel

Compare to Zyrtec® active ingredient

Allergy Relief

Cetirizine Hydrochloride Tablets, 10 mg

Antihistamine

Indoor & Outdoor Allergies

Original Prescription Strength

24 Hour Relief of:

Sneezing

Runny Nose

Itchy, Watery Eyes

Itchy Throat or Nose

24 Hour

actual size

120 TABLETS



Tablets, 10 mg

Antihistamine

24 Hour Relief of:

Runny Nose
 Itchy, Watery Eyes
 Itchy Throat or Nose

Sneezing

Indoor & Outdoor Allergles

Original Prescription Strength

Drug Facts

Active Ingredient (In each tablet)

Purpose .Antihistamine

Drug Facts (continued) reaction to this product occurs. Seek medica help right away.

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Indoor & Outdoor All ergles

Cetirizine Hydrochloride Allergy Relief

Antihistamine Tablets, 10 mg

H-E-B

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Other Information

4H276 %J C5

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Drug Facts (continued)

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

MADE WITH PRIDE & CARE FOR H-E-B® SAN ANTONIO, TX 78204

GLUTEN FREE



Cetirizine HCl 10 ma

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120 TABLETS

size



ALLERGY RELIEF

cetirizine hydrochloride tablet, film coated

Product Information

HUMAN OTC DRUG NDC:37808-583 Product Type Item Code (Source)

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
	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: 3WJQ0SDW1A)			
PO VIDONE, UNSPECIFIED (UNII: FZ989GH94E)			
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)			
TRIACETIN (UNII: XHX3C3X673)			

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

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-583-06	1 in 1 CARTON	01/26/2018	
1		70 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:37808-583-58	1 in 1 CARTON	01/26/2018	
2		40 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:37808-583-39	1 in 1 CARTON	01/26/2018	
3		30 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:37808-583-66	14 in 1 CARTON	0 1/26/20 18	
4		$1\ \text{in}\ 1\ \text{BLISTER}$ PACK; Type 0: Not a Combination Product		
5	NDC:37808-583-76	1 in 1 CARTON	0 1/26/20 18	
5		120 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	ANDA078336	0 1/26/20 18	

Labeler - HEB (007924756)

Revised: 9/2020 HE B