

# **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 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 20-25°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<sup>®</sup> 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



# ALLERGY RELIEF

cetirizine hydrochloride tablet, film coated

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:37808-583
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CETIRIZINE HYDROCHLORIDE</b> (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	10 mg

## Inactive Ingredients

Ingredient Name	Strength
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>FD&amp;C BLUE NO. 1 ALUMINUM LAKE</b> (UNII: J9EQA3S2JM)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I3O)	
<b>POLYDEXTROSE</b> (UNII: VH2XOU12IE)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POVIDONE, UNSPECIFIED</b> (UNII: FZ989GH94E)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>TRIACETIN</b> (UNII: XHX3C3X673)	

## Product Characteristics

<b>Color</b>	WHITE	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	10mm
<b>Flavor</b>		<b>Imprint Code</b>	4H2
<b>Contains</b>			

## 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	10/31/2021
2		40 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:37808-583-39	1 in 1 CARTON	01/26/2018	
		20 in 1 BOTTLE; Type 0: Not a Combination Product		

3		50 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:37808-583-66	14 in 1 CARTON	01/26/2018	
4		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:37808-583-76	1 in 1 CARTON	01/26/2018	
5		120 in 1 BOTTLE; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
<b>Marketing Category</b>		<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ANDA		ANDA078336	01/26/2018	

**Labeler -** H E B (007924756)

Revised: 10/2025

H E B