

# **ALL DAY ALLERGY RELIEF D- cetirizine hcl, pseudoephedrine hcl tablet, extended release H E B**

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**HEB All Day Allergy Relief-D Drug Facts**

## **Active ingredient (in each extended release tablet)**

Cetirizine HCl 5 mg

Pseudoephedrine HCl 120 mg

## **Purpose**

Antihistamine

Nasal Decongestant

## **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
- nasal congestion
- reduces swelling of nasal passages
- temporarily relieves sinus congestion and pressure
- temporarily restores freer breathing through the nose

## **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.
- if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

### **Ask a doctor before use if you have**

- heart disease
- thyroid disease
- diabetes
- glaucoma
- high blood pressure
- trouble urinating due to an enlarged prostate gland
- 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**

- do not use more than directed
- 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.
- you get nervous, dizzy, or sleepless
- symptoms do not improve within 7 days or are accompanied by fever

**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.

**Directions**

- do not break or chew tablet; swallow tablet whole

adults and children 12 years and over	take 1 tablet every 12 hours; do not take more than 2 tablets in 24 hours.
adults 65 years and over	ask a doctor
children under 12 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)

**Inactive ingredients**

colloidal silicon dioxide, hydroxypropyl cellulose, hypromellose, lactose monohydrate, magnesium stearate, microcrystalline cellulose, talc, yellow iron oxide

**Questions or comments?**

1-800-719-9260

## **Principal Display Panel**

Compare to Zyrtec-D® active ingredients

ALL DAY ALLERGY RELIEF-D

Cetirizine Hydrochloride & Pseudoephedrine Hydrochloride

Extended Release Tablets, 5 mg/120 mg

Antihistamine/Nasal Decongestant

Allergy & Congestion

Original Prescription Strength

Indoor & Outdoor Allergies

ACTUAL SIZE

12 Hour Relief of:

Sneezing

Runny Nose

Sinus Pressure

Itchy, Watery Eyes

Itchy Throat or Nose

Nasal Congestion



**Drug Facts (continued)**  
 (Questions or comments? 1-800-719-9260)  
 Made in India  
 HealthCare, division of McNeil-PC, Inc., distributor of Zyrtec-D®  
 SAN ANTONIO, TX 78204  
 1204



**Drug Facts (continued)**

**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  
 ■ if you are now taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

**Ask a doctor before use if you have**  
 ■ heart disease ■ thyroid disease ■ diabetes ■ glaucoma ■ high blood pressure ■ trouble urinating due to an enlarged prostate gland ■ 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**  
 ■ do not use more than directed  
 ■ 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.

**Drug Facts (continued)**

**Directions**  
 ■ do not break or chew tablet; swallow tablet whole  
 adults and children take 1 tablet every 12 hours, do not take more than 2 tablets in 24 hours.  
 adults 65 years and over ask a doctor  
 children under 12 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)

**Inactive ingredients**  
 coltal silicon dioxide, hydroxypropyl cellulose, hydroxypropyl cellulose, talc, yellow iron oxide, miconcristalline cellulose, lactose monohydrate, magnesium stearate.

**CODE AREA**

: 17962 JF C2

**Drug Facts**  
**Active ingredients**  
 (in each extended release tablet)  
 Purpose  
 Cetirizine HCl 5 mg ..... Antihistamine  
 Pseudoephedrine HCl 120 mg ... Nasal Decongestant

**Uses**  
 ■ temporarily relieves these symptoms due to hay fever or other upper respiratory allergies

**Drug Facts (continued)**

**Warnings**  
 ■ you get nervous, dizzy, or sleepless  
 ■ symptoms do not improve within 7 days or are accompanied by fever  
 ■ if pregnant or 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.



**GUARANTEE**  
 We believe the high quality of this H-E-B product makes it an outstanding value. We hope you'll agree. If not, we'll cheerfully refund your money. Thanks for shopping with us.

**DO NOT USE IF BLISTER UNIT IS BROKEN OR TORN**



Compare to Zyrtec-D® active ingredients\*

NDC 37808-176-62



**ALL DAY ALLERGY RELIEF-D**

**Cetirizine Hydrochloride & Pseudoephedrine Hydrochloride Extended Release Tablets, 5 mg/120 mg**

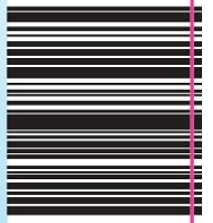
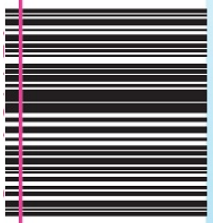
**Antihistamine/Nasal Decongestant**

**Allergy & Congestion**

Original Prescription Strength  
 Indoor & Outdoor Allergies

**12 Hour Relief of:**

ACTUAL SIZE



- Sneezing ■ Runny Nose ■ Sinus Pressure
- Itchy, Watery Eyes ■ Itchy Throat or Nose
- Nasal Congestion

24 EXTENDED  
RELEASE TABLETS

## ALL DAY ALLERGY RELIEF D

cetirizine hcl, pseudoephedrine hcl tablet, extended release

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:37808-176
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>CETIRIZINE HYDRO CHLORIDE</b> (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	5 mg
<b>PSEUDOEPHEDRINE HYDRO CHLORIDE</b> (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	120 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>SILICON DIO XIDE</b> (UNII: ETJ7Z6XBU4)	
<b>HYPROMELLOSES</b> (UNII: 3NXW29V3WO)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>CELLULOSE, MICROCRYSTALLINE</b> (UNII: OP1R32D61U)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	

### Product Characteristics

<b>Color</b>	WHITE (one side white one side light yellow)	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	12mm
<b>Flavor</b>		<b>Imprint Code</b>	5029;5;120
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-176-53	2 in 1 CARTON	05/02/2008	
1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:37808-176-62	4 in 1 CARTON	07/03/2012	
2		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA077170	05/02/2008	

**Labeler** - HEB (007924756)

Revised: 12/2018

HEB