ALLERGY RELIEF- cetirizine hcl tablet Freds Inc

Cetirizine Hydrochloride Tablets

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

Warning

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 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° 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 1-877-753-3935 Monday-Friday 9AM-5PM EST

Principal Display Panel

*Compare to the active ingredient in Zyrtec® ALLERGY RELIEF

Cetirizine HCI

Antihistamine

10 mg tablets

INDOOR & OUTDOOR ALLERGIES

24 HOUR RELIEF OF:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Throat or Nose

TABLETS

*This product is not manufactured or distributed by McNeil Consumer Healthcare, distributor of Zyrtec ${\rm I\!R}$

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOW ANY SIGNS OF TAMPERING

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

DISTRIBUTED BY: fred's Inc.

4300 NEW GETWELL ROAD,

MEMPHIS, TN 38118

www.fredsinc.com

Product Label



Cetirizine HCl antihistamine 10 mg tablets

INDOOR & OUTDOOR ALLERGIES ON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION

VT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER V, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

ALLERGY RELIEF

*Compare to the active ingredient in Zyrtec®

NDC 55315-806-05

INDOOR & OUTDOOR ALLERGIES

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Pharmacy

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Purpose	Directions	
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consumers or kidney o Other in		Warnings Do not use if you have ever had an alle product or any of its ingredients or to an hydroxyzine.
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FRED'S PHARMACY Allergy Relief

ALLERGY RELIEF							
cetirizine hcl tablet							
Product Information							
Product Type	HUMAN OTC	DRUG	Item Code (Source)		NDC:553	NDC:55315-806	
Route of Administration	ORAL						
Active Ingredient/Active	e Moiety						
Ing	redient Nam	ne		Basis of	Strength	Strength	
CETIRIZINE HYDROCHLORIDE UNII:YO7261ME24)	(UNII: 640047K			CETIRIZ INE HYDROCHLOR	RIDE	10 mg	
Inactive Ingredients							
y	Ingredi	ent Name			S	trength	
SILICON DIOXIDE (UNII: ETJ7Z6)	•						
CROSCARMELLOSE SODIUM (U	JNII: M28OL1HH	48)					
HYPROMELLOSE, UNSPECIFIE	D (UNII: 3NXW29	9V3WO)					
LACTOSE MONOHYDRATE (UNI	I: EWQ57Q8I5X)						
MAGNESIUM STEARATE (UNII: 7							
CELLULOSE, MICROCRYSTALL	-	-					
POLYETHYLENE GLYCOL, UNS		I: 3WJQ0SDW1/	4)				
TITANIUM DIOXIDE (UNII: 15FIX	9V2JP)						
Product Characteristics	5						
Color	white	Score			no score		
Shape	OVAL	Size			9mm		
1							

vor Imprint Code			G;4		
Contains					
ackaging	1				
Packaging Markating Start Markating End					
Item Code	ode Package Description	Marketing Start Date	Marketing End Date		
NDC:55315-806- 05	- ⁸⁰⁶⁻ 5 in 1 CARTON	02/28/2019	02/28/2025		
	1 in 1 BLISTER PACK; Type 0: Not a Combination Product	1			
Marketing Information					
Marketing Category		Marketing Start Date	Marketing End Date		
IDA	ANDA209274	02/28/2019	02/28/2025		
DA	ANDAZU92/4	02/20/2019	02/20/2025		

Labeler - Freds Inc (005866116)

Revised: 10/2022

Freds Inc