CHILDRENS ZYRTEC- cetirizine hydrochloride syrup Johnson & Johnson Consumer Inc.

Children's ZYRTEC ®

Drug Facts

Active ingredient (in each 5 mL)

Cetirizine HCl 5 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

- use only with enclosed dosing cup
- find right dose on chart below
- mL = milliliter

adults and children 6 years and over	5 mL or 10 mL once daily depending upon severity of symptoms; do not take more than 10 mL in 24 hours.
adults 65 years and over	5 mL once daily; do not take more than 5 mL in 24 hours.
children 2 to under 6 years of age	2.5 mL once daily. If needed, dose can be increased to a maximum of 5 mL once daily or 2.5 mL every 12 hours. Do not give more than 5 mL in 24 hours.
children under 2 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)
- do not use if carton is opened or if carton tape or bottle wrap imprinted "SAFETY SEAL®" is broken or missing
- see bottom panel for lot number and expiration date

Inactive ingredients

anhydrous citric acid, flavors, propylene glycol, purified water, sodium benzoate,sorbitol solution, sucralose

Questions?

call 1-800-343-7805 (toll-free) or 215-273-8755 (collect)

PRINCIPAL DISPLAY PANEL

NDC 50580-730-05

Children's ZYRTEC[®] ALLERGY

Cetirizine HCl **1 mg /ml** oral solution antihistamine

Indoor & Outdoor Allergies

24

hour Relief of

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

2 yrs. & older

Grape Syrup

Dye-Free • Sugar-Free

4 fl oz (118 ml) Dosing Cup Included

Childrens Cetirizine HCl **1 mg /ml** oral solu ALLERGY Indoor & Outdoor Allergies

> Off Dye-Free • Sugar-Free





ALLERGY

Free • Sugar-Free

Drug Facts	Dru
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Active ingredient Purpose	🖬 us
(in each 5 mL)	■ fin
Cetirizine HCI 5 mgAntihistamine	ml ml
	adult 6yea
Uses	• • • •
temporarity relieves these symptoms due to hay fever or other upper respiratory allergies;	
Tunny nose	adult
III sneezing	and
■ itchy,watery eyes	_
itching of the nose or throat	child
	6 yea
Warnings	
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If pregnant or breast-feeding:	
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■ if pregnant: ask a heelth 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)	





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CHILDRENS ZYRTEC cetirizine hydrochloride syrup **Product Information** HUMAN OTC DRUG NDC:50580-730 **Product Type** Item Code (Source)

Route of Admi	nistration	ORAL					
Active Ingre	dient/Active	Moiety					
	Ingree	dient Name		Basis of Stre	ength	Strer	ngth
CETIRIZINE HYDROCHLORIDE (UNII: 640047KTOA) (CETIRIZINE - UNII:YO7261ME24)			RIZINE -	CETIRIZ INE HYDROCHLORIDE		5 mg in 5 m	L
Inactive Ing	redients						
J		Ingredient Name			Strength		
ANHYDROUS CIT	RIC ACID (UNII: >	-			•••	<u>9</u>	
	COL (UNII: 6DC90						
WATER (UNII: 059							
	ATE (UNII: OJ245F	E5EU)					
SORBITOL SOLU	TION (UNII: 8KW3	E207O2)					
SUCRALOSE (UN	II: 96K6UQ3ZD4)						
Product Cha	racteristics		_				
Color			Score				
Shape			Size				
Flavor		GRAPE Imprint Code					
Contains							
Packaging							
# Item Code	Pa	ckage Description		Marketing Start Date		eting Date	End
1 NDC:50580- 730-05	1 in 1 CARTON			07/13/2015			
1	118 mL in 1 BOT of Co-Package	118 mL in 1 BOTTLE, PLASTIC; Type 1: Convenience Kit of Co-Package					
2 NDC:50580- 730-06	2 in 1 PACKAGE			07/31/2015			
2	1 in 1 CARTON						
2	118 mL in 1 BOT of Co-Package	TLE, PLASTIC; Type 1: 0					
3 NDC:50580- 730-01	1 in 1 CARTON			01/16/2017			
3	30 mL in 1 BOTT of Co-Package	LE, PLASTIC; Type 1: C					
4 NDC:50580- 730-17	3 in 1 PACKAGE			12/03/2018			
4		TLE; Type 1: Convenier	nce Kit of Co-				
	Package						

12/03/2018

5 NDC:50580-730-18

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2 in 1 PACKAGE

1 in 1 CARTON

118 mL in 1 BOTTLE; Type 1: Convenience Kit of Co-Package

ND	A	NDA022155	06/01/2009	
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Marketing Information				
6		240 mL in 1 BOTTLE, PLASTIC; Type 1: Convenience of Co-Package	Kit	
6	730-19	1 in 1 CARTON	06/16/2020	

Labeler - Johnson & Johnson Consumer Inc. (878046358)

Revised: 2/2024

Johnson & Johnson Consumer Inc.