WAL-ZYR ALL DAY ALLERGY- cetirizine hydrochloride solution WALGREEN COMPANY

Wal-Zyr™ Children's Wal-Zyr

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		
<i>j</i> • • • •	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

- do not use if carton is opened or if imprinted safety seal is broken or missing
- see bottom panel for lot number and expiration date
- store between 20° to 25°C (68° to 77°F)

Inactive ingredients

bubble gum artificial flavor, glacial acetic acid, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium acetate anhydrous, sucralose

Questions?

Call **1-866-923-4914**

DISTRIBUTED BY: WALGREEN CO. 200 WILMOT RD., DEERFIELD, IL 60015

PRINCIPAL DISPLAY PANEL - 120 mL Bottle Carton

Walgreens

Compare to Children's Zyrtec[®] active ingredient^{††}

NDC 0363-2106-08

children's ALLERGY

Wal-ZyrTM

CETIRIZINE HYDROCHLORIDE ORAL SOLUTION 1 mg/mL (ANTIHISTAMINE) ALLERGY

DYE & SUGAR FREE

24-HOUR RELIEF

ALCOHOL FREE

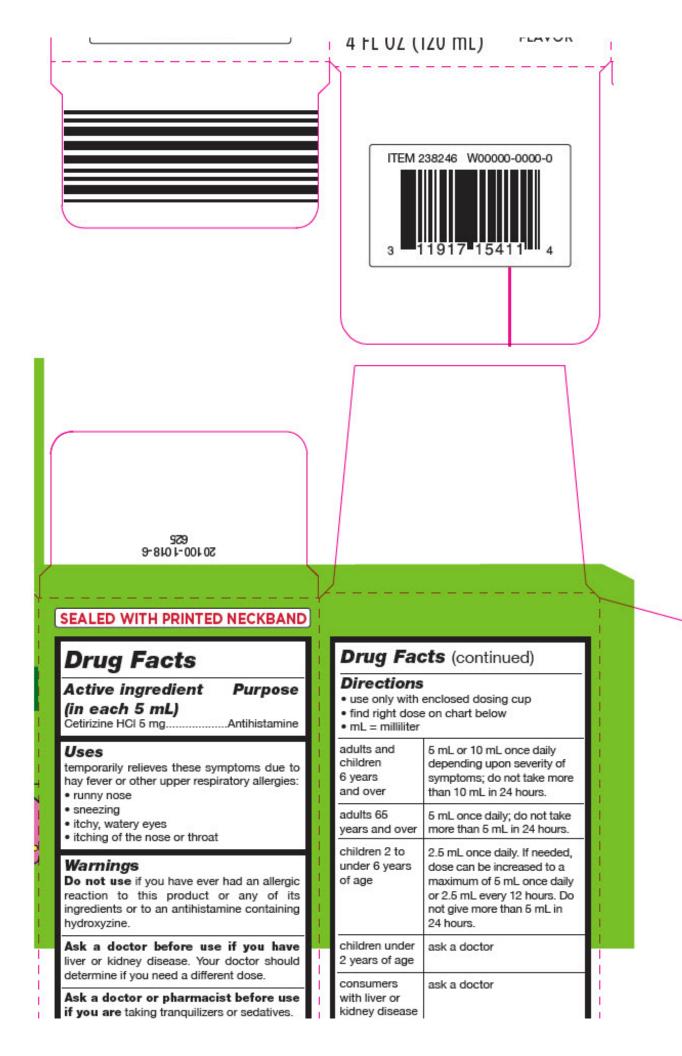
- 24-hour relief of runny nose; sneezing; itchy throat or nose & itchy, watery eyesIndoor & outdoor allergies

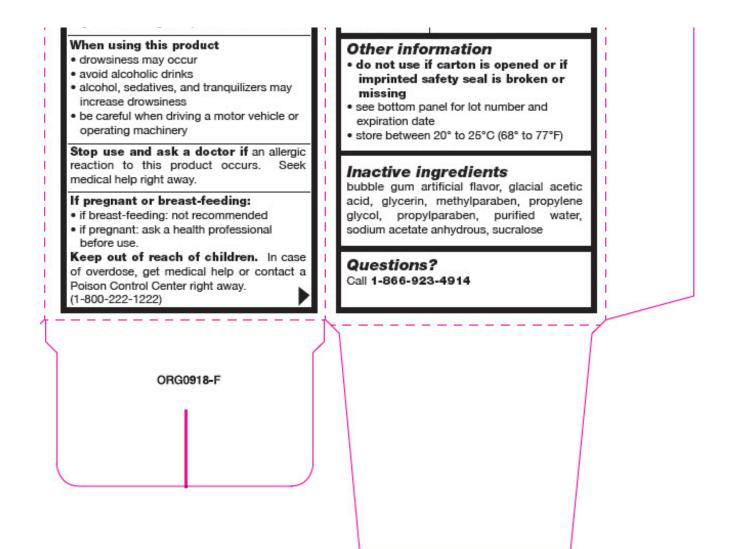
AGES 2 YEARS & OLDER BUBBLE GUM

FLAVOR

4 FL OZ (120 mL)







WAL-ZYR ALL DAY ALLERGY

cetirizine hydrochloride solution

Product Information						
Product T ype	HUMAN OTC DRUG	Item Code (Source)		NDC:036	NDC:0363-2106	
Route of Administration	ORAL					
Active Ingredient/Active Moi	ety					
Ingredient Name Basis of Stree					Strengt	
Cetirizine Hydrochloride (UNII: 640047KTOA) (Cetirizine - UNII:YO7261ME24) Cetirizine Hydrochl					5 mg in 5 m	
Inactive Ingredients						
Ingredient Name				S	Strength	
acetic acid (UNII: Q40Q9N063P)						
glycerin (UNII: PDC6A3C0OX)						
methylparaben (UNII: A2I8C7HI9T)						

рі	propylparaben (UNII: Z8IX2SC1OH)								
w	water (UNII: 059QF0KO0R)								
so	sodium acetate anhydrous (UNII: NVG71ZZ7P0)								
su	sucralose (UNII: 96K6UQ3ZD4)								
P	Product Characteristics								
С	Color YELLOW (colorless to slightly yellow) Sc			Score					
S	Shape			Size					
Flavor BUBBLE GUM Imp			Imprint Code						
С	Contains								
Packaging									
#	Item Code			о р .					
	Item coue	Package Description	Marketing	Start Date	Marketing End	Date			
1	NDC:0363-2106-08	5	Marketing 05/20/2011	Start Date	Marketing End	Date			
1 1		5		Start Date	Marketing End	Date			
1		1 in 1 CARTON 120 mL in 1 BOTTLE; Type 0: Not a Combination Product		Start Date	Marketing End	Date			
1	NDC:0363-2106-08	1 in 1 CARTON 120 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/20/2011	Start Date	Marketing End	Date			
1 2	NDC:0363-2106-08	1 in 1 CARTON 120 mL in 1 BOTTLE; Type 0: Not a Combination Product 1 in 1 CARTON	05/20/2011	Start Date	Marketing End	Date			
1 2	NDC:0363-2106-08	1 in 1 CARTON 120 mL in 1 BOTTLE; Type 0: Not a Combination Product 1 in 1 CARTON	05/20/2011	Start Date	Marketing End	Date			
1 2 2	NDC:0363-2106-08	1 in 1 CARTON 120 mL in 1 BOTTLE; Type 0: Not a Combination Product 1 in 1 CARTON 240 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/20/2011	Start Date	Marketing End	Date			
1 2 2	NDC:0363-2106-08 NDC:0363-2106-01	1 in 1 CARTON 120 mL in 1 BOTTLE; Type 0: Not a Combination Product 1 in 1 CARTON 240 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/20/2011						
1 2 2 M	NDC:0363-2106-08	1 in 1 CARTON 120 mL in 1 BOTTLE; Type 0: Not a Combination Product 1 in 1 CARTON 240 mL in 1 BOTTLE; Type 0: Not a Combination Product Sormation Y Manual Combination Product Y Manual Combination Product	05/20/2011		Marketing End				

Labeler - WALGREEN COMPANY (008965063)

Registrant - Taro Pharmaceuticals U.S.A., Inc. (145186370)

Establishment

Name	Address	ID/FEI	Business Operations
Taro Pharmaceutical Industries, Ltd.		600072078	MANUFACTURE(0363-2106)

Revised: 2/2019

WALGREEN COMPANY