ZYRTEC ALLERGY- cetirizine hydrochloride tablet, orally disintegrating
Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division

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Zyrtec Allergy

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

Tablet melts in mouth. Can be taken with or without water.

	one 10 mg tablet once daily; do not

adults and children 6 years and over	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° to 25°C (68° to 77°F). Avoid high humidity.
- **do not use if carton or blister unit is opened or broken**

Inactive ingredients

amino methacrylate copolymer, anhydrous citric acid, colloidal silicon dioxide, crospovidone, flavors, hydroxypropyl cellulose, magnesium stearate, mannitol, microcrystalline cellulose, sodium bicarbonate, sodium starch glycolate, sucralose

Questions?

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

PRINCIPAL DISPLAY PANEL

Original Prescription Strength
NDC 50580-778-24

ZYRTEC®
Cetirizine HCl orally disintegrating tablets
10mg/antihistamine

Dissolve Tabs™

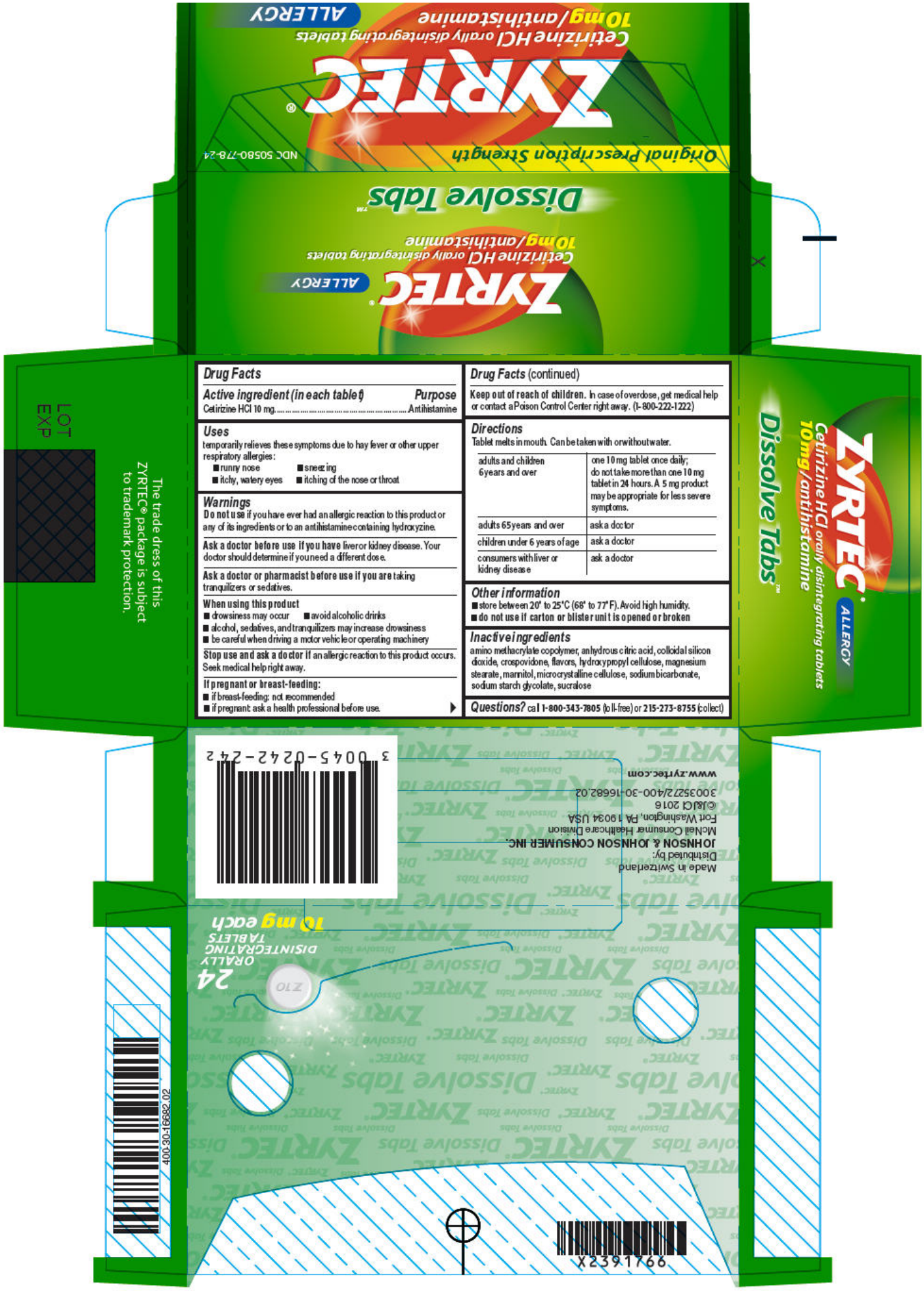
24
HOUR
RELIEF OF

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

ALLERGY
INDOOR + OUTDOOR
ALLERGIES

Melts In
Your Mouth
CITRUS FLAVOR

Actual Size


ZYRTEC®
ALLERGY
Cetirizine HCl orally disintegrating tablets
10mg/antihistamine
Dissolve Tabs™
Original Prescription Strength
NDC 50580-778-24
Drug Facts
Active ingredient (in each tablet) Purpose
Cetirizine HCl 10 mg ..... Antihistamine
Uses
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■ runny nose ■ sneezing
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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
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If pregnant or breast-feeding:
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Drug Facts (continued)
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LOT
EXP
The trade dress of this ZYRTEC® package is subject to trademark protection.
ZYRTEC®
ALLERGY
Cetirizine HCl orally disintegrating tablets
10mg/antihistamine
Dissolve Tabs™
3 0045-0242-242
Made in Switzerland
Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
McNeil Consumer Healthcare Division
Fort Washington, PA 19034 USA
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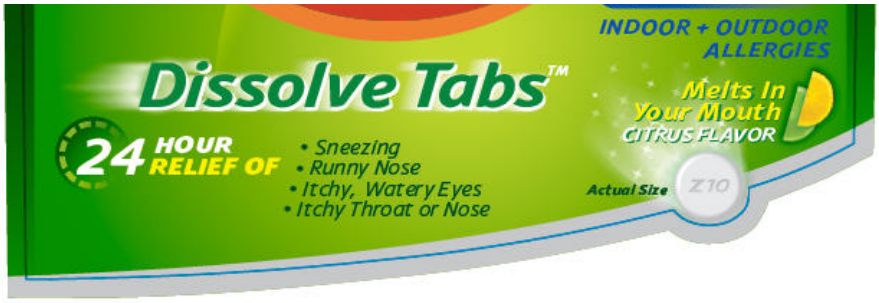
300352724 00-30-16682.02
www.zyrtec.com
24
ORALLY DISINTEGRATING TABLETS
10 mg EACH
Z10
4003016682 02
X2391766



ZYRTEC ALLERGY

cetirizine hydrochloride tablet, orally disintegrating

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50580-778
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Cetirizine Hydrochloride (UNII: 64O047KTOA) (Cetirizine - UNII:YO7261ME24)	Cetirizine Hydrochloride	10 mg

Inactive Ingredients

Ingredient Name	Strength
dimethylaminoethyl methacrylate - butyl methacrylate - methyl methacrylate copolymer (UNII: 905HNO1SIH)	
anhydrous citric acid (UNII: XF417D3PSL)	
silicon dioxide (UNII: ETJ7Z6XBU4)	
crospovidone (15 MPA.S AT 5%) (UNII: 68401960MK)	
hydroxypropyl cellulose (70000 WAMW) (UNII: 66O7AQV0RT)	
magnesium stearate (UNII: 70097M6I30)	
mannitol (UNII: 3OWL53L36A)	
microcrystalline cellulose (UNII: OP1R32D61U)	
sodium bicarbonate (UNII: 8MDF5V39QO)	
sodium starch glycolate type a potato (UNII: 5856J3G2A2)	
sucralose (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color	WHITE (White to Off-white)	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor	CITRUS (citrus-ice)	**Imprint Code**	Z10
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50580-778-12	2 in 1 CARTON	01/20/2014	04/30/2017

1		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:50580-778-24	4 in 1 CARTON	01/20/2014	
2		6 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:50580-778-66	11 in 1 CARTON	01/20/2014	03/31/2017
3		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
NDA	NDA022578	01/20/2014	

Labeler - Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division (878046358)

Revised: 12/2019 Johnson & Johnson Consumer Inc., McNeil Consumer Healthcare Division