# ALLERGY- fexofenadine hydrochloride tablet 7-Eleven

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#### Fexofenadine HCI Tablets USP

## Active ingredient (in each tablet)

Fexofenadine HCI USP, 180 mg

## **Purpose**

**Antihistamine** 

#### Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- itchy, watery eyes
- sneezing
- 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.

# Ask a doctor before use if you have

kidney disease. Your doctor should determine if you need a different dose.

# When using this product

- do not take more than directed
- do not take at the same time as aluminum or magnesium antacids
- do not take with fruit juices (see Directions)

# Stop use and ask doctor if

an allergic reaction to this product occurs. Seek medical help right away.

# If pregnant or breast-feeding,

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**

	take one 180 mg tablet with water once a day; do not take more than 1 tablet in 24 hours
children under 12 years of age	do not use
adults 65 years of age and older	ask a doctor
consumers with kidney disease	ask a doctor

#### Other information

- safety sealed: do not use if carton is opened or if printed foil inner seal on bottle is torn or missing
- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture
- this product meets the requirements of USP Dissolution Test 4

## **Inactive ingredients**

colloidal silicon dioxide, corn starch, croscarmellose sodium, FD&C Red no. 40, hypromellose, iron oxide black, magnesium stearate, mannitol, polyethylene glycol, powder cellulose and titantium dioxide

## **Questions?**

call **1-888-375-3784** 

#### Carton



## **ALLERGY**

fexofenadine hydrochloride tablet

#### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:10202-107(NDC:55111-784)

Route of Administration ORAL

### **Active Ingredient/Active Moiety**

Ingredient Name	<b>Basis of Strength</b>	Strength
Forefore dies Underschlaside (UNIII, 20060D7F711) (FEVOFENADINE	Fovefonadina	

Fexofenadine Hydrochloride (UNII: 2S068B75ZU) (FEXOFENADINE - UNII: E6582LOH6V)

Fexofenadine Hydrochloride Hydrochloride

Inactive Ingredients		
Ingredient Name	Strength	
SILICON DIOXIDE (UNII: ETJ7Z 6XBU4)		
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)		
magnesium stearate (UNII: 70097M6I30)		
mannitol (UNII: 3OWL53L36A)		
POWDERED CELLULOSE (UNII: SMD1X3XO9M)		

FD&C RED NO. 40 (UNII: WZB9127XOA)	
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
polyethylene glycol 400 (UNII: B697894SGQ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics				
Color	PINK	Score	no score	
Shape	OVAL	Size	7mm	
Flavor		Imprint Code	194;R	
Contains				

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:10202-107- 10	2 in 1 CARTON	11/15/2019		
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product			

Marketing Information					
Marketing Application Number or Monograph Marketing Start Marketing End Category Citation Date Date					
ANDA	ANDA076502	11/15/2019			

# **Labeler -** 7-Eleven (007347602)

Revised: 11/2019 7-Eleven