

**ALLERGY RELIEF- fexofenadine hydrochloride tablet**  
**AAA PHARMACEUTICAL, INC.**

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**1192B-RES-2021-0706**

***Drug Facts***

**Active ingredient (in each tablet)**

Fexofenadine HCl 180 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.

**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 a 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 (1-800-222-1222).

**Directions**

adults and children	take one 180 mg tablet with water once a
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12 years of age and over	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

- store between 20-25°C (68-77°F)
- protect from excessive moisture
- retain carton for complete product information and warnings

### Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, lactose monohydrate, macrogol, magnesium stearate, microcrystalline cellulose, pregelatinized starch, titanium dioxide

### Questions or comments?

1-844-705-4384

### PRINCIPAL DISPLAY PANEL

Restore U

NDC 57344-292-04

†COMPARE TO THE ACTIVE INGREDIENT IN ALLEGRE® 24 HOUR

NON-DROWSY

Allergy Relief

Fexofenadine Hydrochloride Tablets, 180 mg / Antihistamine

Indoor / Outdoor Allergies

Relieves:

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

Actual Size

30 TABLETS



## ALLERGY RELIEF

fexofenadine hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:57344-292
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>FEXOFENADINE HYDROCHLORIDE</b> (UNII: 2S068B75Z U) (FEXOFENADINE - UNII:E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	180 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>FERROSFERRIC OXIDE</b> (UNII: XM0M87F357)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX43802MRT)	
<b>HYPROMELLOSE 2910 (15 MPA.S)</b> (UNII: 365FW2JZ0W)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6130)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYETHYLENE GLYCOL 6000</b> (UNII: 30IQX730WE)	

SILICON DIOXIDE (UNII: ETJ7Z6XBU4)

STARCH, CORN (UNII: O8232NY3SJ)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

### Product Characteristics

<b>Color</b>	orange ((PEACH))	<b>Score</b>	no score
<b>Shape</b>	OVAL (Capsule-shaped)	<b>Size</b>	17mm
<b>Flavor</b>		<b>Imprint Code</b>	G6
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57344-292-03	1 in 1 CARTON	07/06/2021	
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:57344-292-04	1 in 1 CARTON	07/06/2021	
2		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:57344-292-06	1 in 1 CARTON	07/06/2021	
3		45 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:57344-292-02	1 in 1 CARTON	07/06/2021	
4		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
5	NDC:57344-292-17	1 in 1 CARTON	07/24/2024	
5		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211075	07/06/2021	

**Labeler** - AAA PHARMACEUTICAL, INC. (181192162)

Revised: 7/2024

AAA PHARMACEUTICAL, INC.