

**ALLERGY RELIEF- fexofenadine hydrochloride tablet**  
**YYBA CORP**

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**ALLERGY RELIEF**

***Drug Facts***

***Active ingredient (in each tablet)***

Fexofenadine HCl 60 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 (1-800-222-1222) right away.

### Directions

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<b>adults and children 12 years of age and over</b>	take one 60 mg tablet with water every 12 hours; do not take more than 2 tablets in 24 hours
<b>children under 12 years of age</b>	do not use
<b>adults 65 years of age and older</b>	ask a doctor
<b>consumers with kidney disease</b>	ask a doctor

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### Other information

- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture
- **each tablet contains:** sodium 2.7 mg
- this product meets the requirements of *USP Dissolution Test 2*

### Inactive ingredients

anhydrous lactose, colloidal silicon dioxide, corn starch, croscarmellose sodium, hypromellose, lactose monohydrate, polyethylene glycol, pregelatinized starch (maize), red iron oxide, stearic acid, titanium dioxide, yellow iron oxide.

### Questions?

call toll-free 1-800-935-6737

### Package Labeling:



<b>Drug Facts (continued)</b>	
<b>Directions</b>	
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children under 12 years of age	do not use
adults 65 years of age and older	ask a doctor
consumers with	ask a doctor

NDC 73581-103-10

Compare to the active ingredient in Allegra® Allergy 12 Hour Tablets\*

60 MG

Gluten Free

**ORIGINAL**  
PRESCRIPTION STRENGTH  
100 TABLETS

DO NOT USE IF PRINTED SEAL UNDER CAP IS MISSING OR DAMAGED

**Drug Facts**

Active ingredient (in each tablet)	Purpose
Fexofenadine HCl 60 mg.....	Antihistamine

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LIFT HERE

PEEL FOR DIRECTIONS      G7024-100-103-0

Kidney disease

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\*This product is not manufactured or distributed by the owner of the registered trademark Allegra® Allergy 12 Hour Tablets



ANTIHISTAMINE

**Allergy Relief**

Made in the USA

Distributed by:  
Wellspring  
Airmont, NY 10952, U.S.A.

Why pay more?  
wellspringmeds.com

## ALLERGY RELIEF

fexofenadine hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:73581-103
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>FEXOFENADINE HYDROCHLORIDE</b> (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	60 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>ANHYDROUS LACTOSE</b> (UNII: 3SY5LH9PMK)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>CORN</b> (UNII: 0N8672707O)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	

**Product Characteristics**

<b>Color</b>	pink	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	12mm
<b>Flavor</b>		<b>Imprint Code</b>	SG;201
<b>Contains</b>			

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:73581-103-10	1 in 1 BOX	06/30/2020	
1		100 in 1 BOTTLE; Type 0: Not a Combination Product		

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
ANDA	ANDA204507	06/30/2020	

**Labeler** - YYBA CORP (006339772)