

**ALLERGY RELIEF- fexofenadine hydrochloride 60 mg tablet**  
**Strategic Sourcing Services, LLC**

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**Fexofenadine HCl Tablets USP**

**Active ingredient(s)**

Fexofenadine HCl USP, 60 mg

**Purpose**

Antihistamine

**Use(s)**

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

adults and children 12 years of age and over	take one 60 mg tablet with water once every 12 hours; do not take more than 2 tablets 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 titanium dioxide

## Questions?

Call **1-888-375-3784**

## PACKAGE LABEL PRINCIPAL DISPLAY PANEL SECTION

1 5 0 0 9 2 4 8 8

1 5 0 0 9 2 4 8 8

**Drug Facts**  
**Active ingredient (in each tablet)** Fexofenadine USP, 60 mg. At that time  
**Purpose** Temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:  
 ■ Runny nose  
 ■ Itchy, watery eyes  
 ■ Sneezing  
 ■ Itching or throbbing of 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 magnesium antacids  
 ■ do not take with fruit juices (see other directions)  
 Stop use and ask a doctor if a 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).

**Drug Facts (continued)**  
**Directions** Adults and children 12 years of age and older: take one (60 mg) tablet with water every 12 hours, do not take more than 2 tablets in 24 hours.  
 Children under 12 years of age: do not use.  
 Adults 65 years of age and older: ask a doctor.  
 Older adults with kidney disease: ask a doctor.

**Other information**  
 ■ Stay sealed; do not use if carton is opened or if individual blister unit is torn or opened.  
 ■ Store between 20°C and 25°C (68°F and 77°F).  
 ■ Protect from excessive moisture.  
 ■ This product meets the requirements of USP Dissolution Test 4.

**Inactive ingredients** croscarmellose sodium, FD-3 colored iron oxide, corn starch, croscarmellose sodium, FD-302 polyethylene glycol powder, colloidal silicon dioxide.  
**Question & Answer** Call 1-800-222-1222 for more information.

TAMPER EVIDENT: INDIVIDUAL BLISTER UNIT SEALED FOR YOUR PROTECTION. DO NOT USE IF INDIVIDUAL BLISTER UNIT IS OPEN OR TORN.  
 IMPORTANT: READ THE DIRECTIONS AND WARNINGS BEFORE USE. KEEP THE CARTON. IT CONTAINS IMPORTANT INFORMATION.

Foster & Thrive™  
 12 HOUR  
**Allergy Relief**  
 Fexofenadine Hydrochloride  
 Tablets USP, 60 mg  
 ANTIHISTAMINE

NDC 70677-1008-1  
 Foster & Thrive™  
 12 HOUR  
**Allergy Relief**  
 Fexofenadine Hydrochloride  
 Tablets USP, 60 mg  
 ANTIHISTAMINE

Non-Drowsy  
 INDOOR/OUTDOOR ALLERGY RELIEF

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



ACTUAL SIZE  
**24 TABLETS**

Foster & Thrive™  
 12 HOUR  
**Allergy Relief**  
 Fexofenadine Hydrochloride  
 Tablets USP, 60 mg  
 ANTIHISTAMINE

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 McNeil Consumer Corporation  
 www.fosterandthrive.com  
 Product of India  
 REV 03/23

\*This product is not manufactured or distributed by  
 Chiltern, Inc. (certified). Scientific Group, distributor of  
 Allergo 12 Hour Tablets, Allergo 5 to 10 mg tablets  
 manufactured by Veranda, LLC.

COMPARE TO THE ACTIVE INGREDIENT IN ALLEGRA® ALLERGY 12 HOUR TABLETS\*



LOT  
 EXP

**ALLERGY RELIEF**  
 fexofenadine hydrochloride 60 mg tablet

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:70677-1008
<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>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>magnesium stearate</b> (UNII: 70097M6I30)	
<b>mannitol</b> (UNII: 3OWL53L36A)	
<b>POWDERED CELLULOSE</b> (UNII: SMD1X3XO9M)	
<b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)	
<b>HYPROMELLOSE 2910 (6 MPA.S)</b> (UNII: 0WZ8WG20P6)	
<b>FERROSFERRIC OXIDE</b> (UNII: XM0M87F357)	
<b>polyethylene glycol 400</b> (UNII: B697894SGQ)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	

## Product Characteristics

<b>Color</b>	PINK	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	5mm
<b>Flavor</b>		<b>Imprint Code</b>	193;R
<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:70677-1008-1	4 in 1 CARTON	05/05/2023	
1		6 in 1 BLISTER PACK; 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	ANDA076502	07/30/2021	

Revised: 11/2024

Strategic Sourcing Services, LLC