

**FEXOFENADINE HCL- fexofenadine hcl tablet, film coated**  
**Camber Consumer Care**

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**FEXOFENADINE HYDROCHLORIDE TABLETS USP, 60mg and 180 mg**

**Allergy**

**ALLERGY**

**Active ingredient (in each film-coated tablet)**

Fexofenadine HCl USP, 60mg and 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 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 (1800-222-1222).

**Directions**

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**FOR 60mg**

adults and children 12 years of age and over	take two 60mg tablets 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
consumers with kidney disease	ask a doctor

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**FOR 180mg**

adults and children 12 years of age and over	take one 180mg 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

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**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
- each tablet contains: sodium 8.2 mg (for 180mg)
- each tablet contains: sodium 2.7 mg (for 60mg)
- this product meets the requirements of USP Dissolution Test 2

**Inactive ingredients**

anhydrous lactose, colloidal silicon dioxide, corn starch, croscarmellose sodium, lactose monohydrate, pregelatinized starch (maize), stearic acid, opadry pink 03B84893 containing hypromellose, polyethylene glycol, red iron oxide, titanium dioxide and yellow iron oxide.

## Questions or comments?

Call 1-888-588-1418

Distributed by:

Camber Consumer Care, Inc.  
Piscataway, NJ 08854, USA,

Made in USA

## Camber Consumer Care - Compare to the active ingredient in Allegra® Allergy 24 Hour Tablets Allergy Relief - 24 HOUR FEXOFENADINE HYDROCHLORIDE TABLETS USP, 180 mg Antihistamine Indoor & Outdoor Allergies



**CAMBER CONSUMER CARE**

**NDC 69230-201-01 Compare to the active ingredient in Allegra® Allergy\***

**Non-Drowsy**

**Fexofenadine Hydrochloride Tablets USP**

**60 mg**


**Antihistamine**

**12 HR Indoor/Outdoor Allergy Relief**

**Sneezing**

**Runny Nose**

# Itchy, Watery Eyes Itchy Nose or Throat

 <p>NDC 69230-201-01 Compare to the active ingredient in Allegra® Allergy*</p> <p><b>Non-Drowsy</b> <b>Fexofenadine Hydrochloride Tablets USP</b> <b>60 mg</b> Antihistamine 12 HR Indoor/Outdoor Allergy Relief - Sneezing - Runny Nose - Itchy, Watery Eyes - Itchy Nose or Throat <b>100 Tablets Allergy</b></p>	<p><b>TAMPER EVIDENT: DO NOT USE IF FOIL SEAL UNDERCAP PRINTED WITH "SEALED FOR YOUR PROTECTION" IS BROKEN OR MISSING</b></p>	<p><b>Drug Facts</b></p>	<p><b>Active ingredient</b> (in each film-coated tablet) Fexofenadine HCl USP, 60 mg ..... Antihistamine</p>	<p><b>Purpose</b> 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</p>	<p><b>Warnings</b> Do not use if you have ever had an allergic reaction to this product or any of its ingredients.</p>	<p><b>Ask a doctor before use if you have</b> kidney disease. Your doctor should determine if you need a different dose.</p>	<p><b>When using this product</b></p> <ul style="list-style-type: none"> <li>■ do not take more than directed</li> <li>■ do not take at the same time as aluminum or magnesium antacids</li> <li>■ do not take with fruit juices (see Directions) ▶</li> </ul>	<p>(Continued On Back Of Label) *This product is not manufactured or distributed by Chattem, Inc., a Sanofi Company, distributor of Allegra® Allergy</p> <p>Distributed by: Camber Consumer Care, Inc. Piscataway, NJ 08854, USA Made in USA</p> <p>Rev. 08/23</p> <p>LOT EXP</p> <p>PEEL HERE</p> <p>N 3 6923020101 5</p> <p>0.2" x 0.8" NO VARNISH</p>
		<p><b>ADHESIVE AREA</b></p> <p><b>Drug Facts</b> (continued) <b>Stop use and ask a doctor if an allergic reaction to this product occurs.</b> Seek medical help right away. <b>If pregnant or breast-feeding,</b> ask a health professional before use. <b>Keep out of reach of children.</b> In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). <b>Directions</b> ■ <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 <b>Other information</b> ■ store between 20° and 25°C (68° to 77°F) ■ protect from excessive moisture ■ each tablet contains sodium 2.7 mg ■ this product meets the requirements of USP Dissolution Test 2 <b>Inactive ingredients</b> anhydrous lactose, colloidal silicon dioxide, corn starch, croscarmellose sodium, lactose monohydrate, pregelatinized starch (maize), stearic acid, opadry pink 03B84893 containing hypromellose, polyethylene glycol, red iron oxide, titanium dioxide and yellow iron oxide <b>Questions?</b> Call 1-888-588-1418</p>						

<p><b>ADHESIVE AREA</b></p>	<p><b>ADHESIVE AREA</b></p>
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NDC 69230-201-05 Compare to the active ingredient in Allegra® Allergy\*

Non-Drowsy

Fexofenadine Hydrochloride Tablets USP

60 mg

Antihistamine

12 HR Indoor/Outdoor Allergy Relief

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

NDC 69230-201-05  
Compare to  
the active ingredient in  
Allegra® Allergy\*

**CAMBER**  
CONSUMER CARE

**Non-Drowsy  
Fexofenadine  
Hydrochloride  
Tablets USP**

**60 mg**

**Antihistamine**

**12 HR  
Indoor/Outdoor Allergy Relief**

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

**500 Tablets Allergy**

TAMPER EVIDENT DO NOT USE IF FOIL SEAL UNDERCAP PRINTED WITH "SEALED FOR YOUR PROTECTION" IS BROKEN OR MISSING

<b>Drug Facts</b>	<p><b>Purpose</b> Antihistamine</p> <p><b>Active ingredient</b> (in each film-coated tablet) Fexofenadine HCl USP, 60 mg</p> <p><b>Uses</b> 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</p> <p><b>Warnings</b> Do not use if you have ever had an allergic reaction to this product or any of its ingredients.</p> <p><b>Ask a doctor before use if you have kidney disease.</b> Your doctor should determine if you need a different dose.</p> <p><b>When using this product</b> ■ 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)</p> <p><b>Stop use and ask a doctor if</b> an allergic reaction to this product occurs. Seek medical help right away.</p> <p><b>If pregnant or breast-feeding, ask a health professional before use. Keep out of reach of children.</b> In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222).</p> <p><b>Directions</b> ■ <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</p> <p><b>Other information</b> ■ store between 20° and 25°C (68° to 77°F) ■ protect from excessive moisture ■ each tablet contains sodium 2.7 mg ■ this product meets the requirements of USP Dissolution Test 2</p> <p><b>Inactive ingredients</b> anhydrous lactose, colloidal silicon dioxide, corn starch, croscarmellose sodium, lactose monohydrate, pregelatinized starch (maize), stearic acid, polyethylene glycol, red iron oxide, titanium dioxide and yellow iron oxide.</p> <p><b>Questions or comments?</b> call 1-888-588-1418</p> <p><small>*This product is not manufactured or distributed by Chiesmi, Inc., a Sanofi Company, distributor of Allegra® Allergy. Distributed by: Cambor Consumer Care, Inc. Piscataway, NJ 08854, USA Made in USA Rev. 09/21</small></p>
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LOT

3 EXP

NO VARNISH

0.55" x 0.82"

6 9 2 3 0 2 0 1 0 5

# FEXOFENADINE HCL

fexofenadine hcl tablet, film coated

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69230-202
<b>Route of Administration</b>	ORAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>FEXOFENADINE HYDROCHLORIDE</b> (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	180 mg

## Inactive Ingredients

Ingredient Name	Strength
<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>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX43802MRT)	

## Product Characteristics

<b>Color</b>	pink	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	17mm
<b>Flavor</b>		<b>Imprint Code</b>	SG;202



**Contains****Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69230-202-30	1 in 1 CARTON	09/16/2015	
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:69230-202-45	1 in 1 CARTON	09/16/2015	
2		45 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:69230-202-60	1 in 1 CARTON	09/16/2015	
3		60 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:69230-202-90	1 in 1 CARTON	09/16/2015	
4		90 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:69230-202-01	1 in 1 CARTON	09/16/2015	
5		100 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:69230-202-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	09/16/2015	
7	NDC:69230-202-11	1000 in 1 BOTTLE; Type 0: Not a Combination Product	09/16/2015	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204507	09/16/2015	

**FEXOFENADINE HCL**

fexofenadine hcl tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:69230-201
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

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

## Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS LACTOSE</b> (UNII: 3SY5LH9PMK)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<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 (Modified Oval)	<b>Size</b>	12mm
<b>Flavor</b>		<b>Imprint Code</b>	SG;201
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69230-201-05	500 in 1 BOTTLE; Type 0: Not a Combination Product	12/08/2021	
2	NDC:69230-201-01	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/08/2021	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204507	12/08/2021	

**Labeler** - Camber Consumer Care (079539968)

Revised: 11/2023

Camber Consumer Care