

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

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

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

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

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 69230-201-05 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

CAMBER CONSUMER CARE
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
500 Tablets Allergy

Drug Facts
Active ingredient (in each film-coated tablet) Fexofenadine HCl USP, 60 mg 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 (1-800-222-1222).
Directions ■ adults and children 12 years of age and over 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 ■ consumers with kidney disease ask a doctor
Other information ■ 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
Inactive ingredients anhydrous lactose, colloidal silicon dioxide, corn starch, croscarmellose sodium, lactose monohydrate, pregelatinized starch (maize), stearic acid, opadry pink, 0.3B84893 containing hypromellose, polyethylene glycol, red iron oxide, titanium dioxide and yellow iron oxide.
Questions or comments? call 1-888-588-1418
This product is not manufactured or distributed by Chatham, Inc., a Sunovion Company, distributor of Allegra® Allergy.
Distributed by: Camber Consumer Care, Inc. Piscataway, NJ 08854, USA
Made in USA
Rev. 09/21
0.55" x 0.82" NO VARNISH
LOT 69230201051 3 EXP

FEXOFENADINE HCL

fexofenadine hcl tablet, film coated

Product Information

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

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
TITANIUM DIOXIDE (UNII: 15F1X9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	

Product Characteristics

Color	pink	Score	no score
Shape	CAPSULE	Size	17mm
Flavor		Imprint Code	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

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

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	60 mg
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Inactive Ingredients	
Ingredient Name	Strength
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	

Product Characteristics			
Color	pink	Score	no score
Shape	OVAL (Modified Oval)	Size	12mm
Flavor		Imprint Code	SG;201
Contains			

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	

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)