

FEXOFENADINE HCL- fexofenadine hcl tablet, film coated
AKRON PHARMA INC

ALLERGY

Active ingredient (in each film-coated tablet)

Fexofenadine HCl USP 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 right away.

Directions (for 60mg)

adults and children 12 years of age and over	take one 60mg 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° and 77°F)
- protect from excessive moisture
- **each tablet contains:**sodium 2.7mg(for 60 mg)
- this product meets the requirements of USP *Dissolution Test 2*
- **Tamper Evident:**Do not use if imprinted inner safety seal is torn or missing

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 and yellow iron oxide.

Questions or comments?

Call toll-free 1-877-225-6999

Distributed by:

Akron Pharma, Inc.

Fairfield, NJ 07004

www.akronpharma.com

NDC 71399-8659-1

*Compare to the active ingredient in Allegra® Allergy

ORIGINAL PRESCRIPTION STRENGTH, NON-DROWSY

Allergy Relief

Fexofenadine Hydrochloride

Tablets USP, 60 mg

Antihistamine

INDOOR AND OUTDOOR ALLERGIES

12 Hour Relief of:

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

12 Hr

100 Tablets
60 mg Each

Drug Facts	
Active Ingredient (in each film-coated tablet) Fexofenadine HCl USP, 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.	

PEEL HERE

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No Varnish

Akron Pharma
Distributed by:
Akron Pharma, Inc.
Fairfield, NJ 07004
www.akronpharma.com
Made in USA
Rev:06/24

*This Product is not manufactured or distributed by Chatterm, Inc. (part of the Sanofi Group), distributor of Allegra® Allergy tablets. Allegra® is a registered trademark of Aventisub II Inc.

4.50"

Drug Facts (Continued)	
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 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° and 77°F) ■ protect from excessive moisture ■ each tablet contains: sodium 2.7 mg ■ this product meets the requirements of USP Dissolution Test 2 ■ Tamper Evident: Do not use if imprinted inner safety seal is torn or missing.	
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NDC 71399-8659-5

*Compare to the active ingredient in Allegra® Allergy

ORIGINAL PRESCRIPTION STRENGTH, NON-DROWSY

Allergy Relief

Fexofenadine Hydrochloride

Tablets USP, 60 mg

Antihistamine

INDOOR AND OUTDOOR ALLERGIES

12 Hour Relief of:

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

12 Hr

500 Tablets
60 mg Each

Drug Facts	
Active Ingredient (in each film-coated tablet) Fexofenadine HCl USP, 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.	
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Questions or comments? Call toll-free 1-877-225-6999	

0.632" X 1.24"

N 3 71399 86595 3

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FEXOFENADINE HCL

fexofenadine hcl tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC: 71399-8659
Route of Administration	ORAL		
Active Ingredient/Active Moiety			

Ingredient Name		Basis of Strength	Strength	
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V)		FEXOFENADINE HYDROCHLORIDE	60 mg	
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	Size	12mm	
Flavor		Imprint Code	SG;201	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-8659-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	11/01/2021	
2	NDC:71399-8659-5	500 in 1 BOTTLE; Type 0: Not a Combination Product	02/02/2024	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA204507	12/26/2014		

Labeler - AKRON PHARMA INC (067878881)

Revised: 1/2026

AKRON PHARMA INC