

FEXOFENADINE HCL- fexofenadine hcl tablet, film coated
CVS Pharmacy

FEXOFENADINE HYDROCHLORIDE TABLETS USP, 180 mg

ALLERGY

Active ingredient (in each film-coated tablet)

Fexofenadine HCl USP, 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.

Directions

adults and children 12 years of age and older take one 180mg tablet with water once a day; do not take more than 1

age and over	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 individual blister units are torn or opened.
- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture
- each tablet contains: sodium 8.2 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 03B84893 containing hypromellose, polyethylene glycol, red iron oxide, titanium dioxide and yellow iron oxide.

Questions or comments?

Call 1-888-588-1418

CVS Pharmacy, Inc.

One CVS Drive

Woonsocket, RI 02895

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CVS.com® 1-800-SHOP CVS

V-32583

CVS Health-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

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.

5 TABLETS 180 mg EACH

Non-Drowsy

- Sneezing
- Runny nose
- Itchy, watery eyes
- Itchy nose or throat

24 Hour Relief of:

Antihistamine

TABLETS USP, 180 mg

FEXOFENADINE HYDROCHLORIDE

Allergy Relief

Original Prescription Strength

CVS Health

Indoor & Outdoor Allergies

Compare to the active ingredient in Allegra[®] Allergy 24 Hour Tablets[®]



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Drug Facts

Active ingredient (in each film-coated tablet)	Purpose
Fexofenadine HCl USP, 180 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.

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Other information ■ safety sealed; do not use if carton is opened or if individual blister units are torn or opened. ■ store between 20° and 25°C (68° and 77°F) ■ protect from excessive moisture ■ each tablet contains: sodium 8.2 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 08B04893 containing hypromellose, polyethylene glycol, red iron oxide, titanium dioxide and yellow iron oxide

Questions? Call 1-888-588-1418

691007

700160

23020201



Actual Size

LOT
 EXP

FEXOFENADINE HCL

fexofenadine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-979
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FEXO FENADINE HYDRO CHLORIDE (UNII: 2S068B75ZU) (FEXO FENADINE - UNII:E6582LOH6V)	FEXO FENADINE HYDROCHLORIDE	180 mg

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
COLLOIDAL SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	

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:59779-979-81	1 in 1 CARTON		
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:59779-979-83	3 in 1 CARTON		
2		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:59779-979-30	30 in 1 CARTON; Type 0: Not a Combination Product		
4	NDC:59779-979-45	45 in 1 CARTON; Type 0: Not a Combination Product		
5	NDC:59779-979-73	60 in 1 CARTON; Type 0: Not a Combination Product		
6	NDC:59779-979-90	90 in 1 CARTON; Type 0: Not a Combination Product		
7	NDC:59779-979-53	180 in 1 CARTON; Type 0: Not a Combination Product		

Marketing Information

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

Labeler - CVS Pharmacy (062312574)

Revised: 10/2015

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