CABINET ALLERGY RELIEF FEXOFENADINE- fexofenadine hydrochloride tablet Cabinet Health P.B.C.

Cabinet Non Drowsy Allergy Relief, Fexofenadine

Drug Facts

Active ingredient (in each 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. (1-800-222-1222)

Directions

adults and children 12 years of age and over	take one 180 mg 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 imprinted foil under bottle cap is opened or torn
- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture

Inactive ingredients

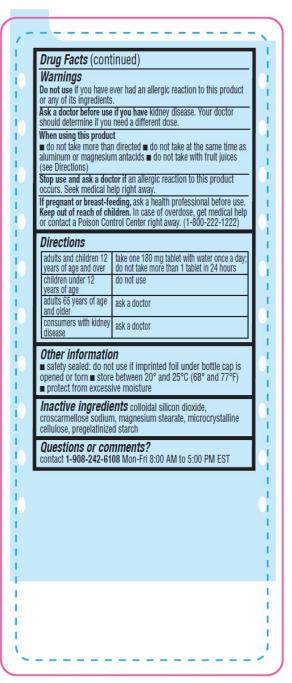
colloidal silicon dioxide, croscarmellose sodium, magnesium stearate, microcrystalline cellulose, pregelatinized starch

Questions or comments?

contact 1-908-242-6108Mon-Fri 8:00 AM to 5:00 PM EST

Package Labeling:





CABINET ALLERGY RELIEF FEXOFENADINE

fexofenadine hydrochloride tablet

	Product Information			
	Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82725-3002
l	Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII: F6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	180 mg

Inactive Ingredients		
Ingredient Name	Strength	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		

Product Characteristics			
Color	pink	Score	no score
Shape	CAPSULE	Size	17mm
Flavor		Imprint Code	G6
Contains			

	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:82725-3002-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2025	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211075	09/01/2025	

Labeler - Cabinet Health P.B.C. (117102391)

Registrant - Cabinet Health P.B.C. (117102391)

Revised: 10/2025 Cabinet Health P.B.C.