

**GOODMEDS NON DROWSY ALLERGY RELIEF- fexofenadine  
hydrochloride tablet  
Cabinet Health P.B.C.**

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**Goodmeds Non Drowsy Allergy Relief**

***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**

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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

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## **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, hypromellose, light liquid paraffin, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, pregelatinized starch, red iron oxide, sodium starch glycolate, talc, titanium dioxide, yellow iron oxide

## **Questions or comments?**

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

## **Package Labeling:**

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**Drug Facts (continued)**

**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**

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**goodmeds** Compare to the active ingredient in ALLEGRA® Allergy 24 Hour†

**NON-DROWSY**

# Allergy relief

Fexofenadine HCl, USP 180 mg, Antihistamine  
For indoor and outdoor allergies

24 hour relief of:

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

120 Tablets

actual size 

NDC 82725-4202-1

**TAMPER EVIDENT: DO NOT USE IF IMPRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING**

**IMPORTANT: READ THE DIRECTIONS AND WARNINGS BEFORE USE**

**Drug Facts**

Active ingredient (in each 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

(continued on back of label)

† This product is not manufactured or distributed by Chattem, Inc., the distributor of Allegra® Allergy 24 Hour. Made in India

Distributed by: Cabinet Health P.B.C. Brooklyn, NY 11222  
LOT: EXP: ORG 01/26

ITEM# GM-4202-1

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**GOODMEDS NON DROWSY ALLERGY RELIEF**

fexofenadine hydrochloride tablet

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:82725-4202
<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>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>LIGHT MINERAL OIL</b> (UNII: N6K5787QVP)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>SODIUM STARCH GLYCOLATE TYPE A</b> (UNII: H8AV0SQX4D)	
<b>TALC</b> (UNII: 7SEV7J4R1U)	
<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>	CAPSULE	<b>Size</b>	18mm
<b>Flavor</b>		<b>Imprint Code</b>	j;44
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82725-4202-1	120 in 1 BOTTLE; Type 0: Not a Combination Product	02/10/2026	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204097	02/10/2026	

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

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

