

**ALLEGRA HIVES 24HR- fexofenadine hydrochloride tablet, film coated  
Chattem, Inc.**

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**Allegra Hives 24HR**

**Allegra Hives 24HR tablet 180 mg**

***Drug Facts***

***Active ingredient (in each tablet)***

Fexofenadine HCl 180 mg

***Purpose***

Antihistamine

***Uses***

reduces hives and relieves itching due to hives (urticaria). This product will not prevent hives or an allergic skin reaction from occurring.

***Warnings***

**Severe Allergy Warning:** Get emergency help **immediately** if you have hives along with any of the following symptoms:

- trouble swallowing ■ dizziness or loss of consciousness ■ swelling of tongue
- swelling in or around mouth ■ trouble speaking ■ wheezing or problems breathing ■ drooling

These symptoms may be signs of anaphylactic shock. This condition can be life threatening if not treated by a health professional **immediately**. Symptoms of anaphylactic shock may occur when hives first appear or up to a few hours later.

**Not a Substitute for Epinephrine.** If your doctor has prescribed an epinephrine injector for “anaphylaxis” or severe allergy symptoms that could occur with your hives, never use this product as a substitute for the epinephrine injector. If you have been prescribed an epinephrine injector, you should carry it with you at all times.

**Do not use**

■ to **prevent** hives from any known cause such as:

■ foods ■ insect stings ■ medicines ■ latex or rubber gloves because this product will not stop hives from occurring. Avoiding the cause of your hives is the only way to prevent them. Hives can sometimes be serious. If you do not know the cause of your hives, see your doctor for a medical exam. Your doctor may be able to help you find a

cause.

- 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.
- hives that are an unusual color, look bruised or blistered
- hives that do not itch

**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.
- symptoms do not improve after 3 days of treatment
- the hives have lasted more than 6 weeks

**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 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**

- (if blister) safety sealed: do not use if carton is opened or if individual blister units are torn or opened
- (on the carton if bottle) safety sealed: do not use if carton is opened or if printed foil

inner seal on bottle is torn or missing

■ (if bottle) safety sealed: do not use if carton was 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

### Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide blends, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch, titanium dioxide

### Questions or comments?

call toll-free 1-800-633-1610

### PRINCIPAL DISPLAY PANEL

Allegra  
Hives  
fexofenadine HCl tablet  
180 mg/antihistamine  
24 HR  
30 TABLETS



# ALLEGRA HIVES 24HR

fexofenadine hydrochloride tablet, film coated

## Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:41167-4126
<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>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<b>BROWN IRON OXIDE</b> (UNII: 1N032N7MFO)	
<b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)	
<b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>POVIDONE</b> (UNII: FZ989GH94E)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

## Product Characteristics

<b>Color</b>	orange (peach)	<b>Score</b>	no score
<b>Shape</b>	OVAL	<b>Size</b>	17mm
<b>Flavor</b>		<b>Imprint Code</b>	018;E
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41167-4126-8	1 in 1 CARTON	06/06/2022	
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:41167-4126-3	1 in 1 CARTON	01/31/2024	
2		15 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:41167-4126-4	1 in 1 CARTON	06/06/2022	
3		30 in 1 BOTTLE; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA020872	06/06/2022	

**Labeler** - Chattem, Inc. (003336013)

Revised: 1/2026

Chattem, Inc.