

**FEXOFENADINE HYDROCHLORIDE- fexofenadine hydrochloride tablet**  
**NUVICARE LLC**

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**Allergy Relief Fexofenadine Hydrochloride Tablets 180 mg**

**Drug Facts**

**Active ingredient (in each tablet)**

Fexofenadine HCl 180 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 (1-800-222-1222) right away.

**Directions**

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adults and children 12 years	take one 180 mg tablet with water once a day; do not take
of age and over	more than 1 tablet in 24 hours
children under 12 years of	do not use

age  
adults 65 years of age and  
older  
consumers with kidney  
disease

do not use  
ask a doctor  
ask a doctor

**Other information** □ store between 20° and 25°C (68° and 77°F) □ protect from excessive moisture

**Inactive ingredients:** colloidal silicone dioxide, croscarmellose sodium, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol/macrogol, povidone, pregelatinized starch, red iron oxide, silica, titanium dioxide, yellow iron oxide.

Call 1(718) 337-8733 or visit: support@nuvicare.com





## FEXOFENADINE HYDROCHLORIDE

fexofenadine hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:84324-008
<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>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	
<b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)	
<b>CROSCARMELLOSE SODIUM</b> (UNII: M28OL1HH48)	
<b>HYPROMELLOSE, UNSPECIFIED</b> (UNII: 3NXW29V3WO)	
<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>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

### Product Characteristics

<b>Color</b>	orange	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	18mm
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84324-008-02	4 in 1 BLISTER PACK; Type 0: Not a Combination Product	06/09/2025	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210137	06/09/2025	

**Labeler** - NUVICARE LLC (119257565)

**Registrant** - NUVICARE LLC (119257565)