

**FEXOFENADINE HYDROCHLORIDE- fexofenadine hydrochloride tablet**  
**Spirit Pharmaceuticals LLC**

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**Valumed**

**FEXOFENADINE**

**HYDROCHLORIDE**

**TABLETS USP**

**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. You may report side effects to FDA at **1-800-FDA-1088**.

**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

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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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## Drug Facts (continued)

**Other information** □ safety sealed: do not use 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 silicone dioxide, croscarmellose sodium, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol/macrogol, povidone, pregelatinized starch, red iron oxide, silica, titanium dioxide, yellow iron oxide.

**Questions or comments?** call **+1-888-333-9792**

Distributed by:  
Spirit Pharmaceuticals LLC,  
Ronkonkoma, NY 11779

ORG 03/19

Manufactured by:  
Unique Pharmaceutical  
Laboratories  
(A Div. of J. B. Chemicals &  
Pharmaceuticals Ltd.)  
Mumbai 400 030, India  
Mfg. Lic. No.: G/1430

**PRINCIPAL DISPLAY PANEL**

Valumed<sup>TM</sup>

NDC 68210-0122-1

non-drowsy

FEXOFENADINE

HYDROCHLORIDE

TABLETS USP

180 mg

antihistamine

24 Hour

indoor & outdoor allergy relief

- sneezing • runny nose
- itchy, watery eyes
- itchy nose or throat

100 tablets

Actual Size

**ValuMeds**  
NDC 68210-0122-1  
non-drowsy  
**FEXOFENADINE  
HYDROCHLORIDE  
TABLETS USP**  
180 mg  
antihistamine

24 Hour  
indoor & outdoor allergy relief  
• sneezing • runny nose  
• itchy, watery eyes  
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100 tablets

Actual Size

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
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consumers with kidney disease	ask a doctor

**Drug Facts**  
**Active ingredient (in each tablet)**  
Fexofenadine HCl 180 mg  
**Purpose**  
Antihistamine  
**Use 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. You may report side effects to FDA at 1-800-FDA-1088. 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**

**Other information** • safety sealed; do not use 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 silicone dioxide, croscarmellose sodium, hypromellose, magnesium stearate, microcrystalline cellulose, polyethylene glycol/macrogol, polydioxanone, pregelatinized starch, red iron oxide, silica, titanium dioxide, yellow iron oxide.

**Questions or comments?** call 1-888-333-8792

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ORG 03/19

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Lot. No. :  
Exp. Date :

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## FEXOFENADINE HYDROCHLORIDE

fexofenadine hydrochloride tablet

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:68210-0122
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name		Basis of Strength	Strength	
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V)		FEXOFENADINE HYDROCHLORIDE	180 mg	
<b>Inactive Ingredients</b>				
Ingredient Name		Strength		
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)				
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)				
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)				
HYPROMELLOSE 2910 (15 MPA.S) (UNII: 36SFW2JZ0W)				
HYPROMELLOSE 2910 (5 MPA.S) (UNII: R75537T0T4)				
MAGNESIUM STEARATE (UNII: 70097M6I30)				
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)				
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)				
POVIDONE (UNII: FZ989GH94E)				
STARCH, CORN (UNII: O8232NY3SJ)				
FERRIC OXIDE RED (UNII: 1K09F3G675)				
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)				
<b>Product Characteristics</b>				
Color	orange	Score	no score	
Shape	CAPSULE	Size	17mm	
Flavor		Imprint Code	180	
Contains				
<b>Packaging</b>				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68210-0122-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	03/22/2019	
2	NDC:68210-0122-3	1 in 1 CARTON	09/18/2020	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
<b>Marketing Information</b>				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA210137	03/21/2019		

**Labeler** - Spirit Pharmaceuticals LLC (179621011)