

**FEXOFENADINE HYDROCHLORIDE- fexofenadine hcl tablet**  
**Northwind Pharmaceuticals, LLC**

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**Active ingredient (in each film-coated tablet)**

Fexofenadine HCL 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

- **each tablet contain:** sodium 8 mg
- store between 20 to 25°C (68 to 77°F)
- protect from excessive moisture
- this product meets the requirements of USP Dissolution Test 2

## Inactive ingredients

anhydrous lactose, colloidal silicon dioxide, corn starch, croscarmellose sodium, hypromellose, lactose monohydrate, polyethylene glycol, pregelatinized starch (maize), red iron oxide, stearic acid, titanium dioxide, and yellow iron oxide

## Questions or comments?

Call: **1-800-616-2471**

## Principal Display Panel

COMPARE TO the active ingredient in ALLEGRA® ALLERGY 24 HOUR\*

## **FEXOFENADINE HYDROCHLORIDE USP 180 mg**

### **Antihistamine**

ALLERGY RELIEF

24-HOUR Relief of:

- sneezing
- Runny nose
- Itchy, watery eyes
- Itchy nose or throat

Indoor and Outdoor

Allergy Relief

Non-Drowsy

Capsule Shaped Tablets

\*This product is not manufactured or distributed by Chattem Inc., distributor of Allegra® Allergy 24 hour

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

Distributed by:

**MAJOR® PHARMACEUTICALS**

17177 N Laurel Park Drive, Suite 233

Livonia, MI 48152

**Product Label**

**NDC: 51655-231-26**

**NDC: 51655-231-26**  
**Fexofenadine Hydrochloride USP, 180 mg Antihistamine 90 Tablets**

Dosage: See package insert  
 Store at 20° - 25°C (68° - 77°F) (See USP Controlled Room Temperature)


Keep out of the reach of children.  
 Store in original container.  
 Protect from excessive moisture. 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

LCN#: 00  
 Rev. A 12/22

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. Each tablet contain: sodium 8 mg. This product meets the requirements of USP. Dissolution test 2. Active Ingredient (in each film-coated tablet) Fexofenadine HCl USP 180mg.

Repackaged From: 0904-7050-XX  
 Major Pharmaceuticals, Lot 0000000000

Repackaged By: Northwind Pharmaceuticals  
 Indianapolis, IN 46203  
 GTIN: 00351655231265  
 S/N: 000000000000000  
 EXP: 00/00/0000  
 LOT: 0000000000



<b>FEXOFENADINE HYDROCHLORIDE</b>			
fexofenadine hcl tablet			
Product Information			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:51655-231(NDC:0904-7050)
<b>Route of Administration</b>	ORAL		
Active Ingredient/Active Moiety			
<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>	
FEXOFENADINE HYDROCHLORIDE (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>HYPROMELLOSES</b> (UNII: 3NXW29V3WO)	
<b>FERRIC OXIDE RED</b> (UNII: 1K09F3G675)	
<b>FERRIC OXIDE YELLOW</b> (UNII: EX438O2MRT)	
<b>ANHYDROUS LACTOSE</b> (UNII: 3SY5LH9PMK)	
<b>STARCH, CORN</b> (UNII: O8232NY3SJ)	
<b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A)	
<b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)	

### Product Characteristics

<b>Color</b>	pink	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	17mm
<b>Flavor</b>		<b>Imprint Code</b>	SG;202
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51655-231-52	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/16/2022	
2	NDC:51655-231-26	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/08/2022	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204507	11/16/2022	

**Labeler** - Northwind Pharmaceuticals, LLC (036986393)

**Registrant** - Northwind Pharmaceuticals, LLC (036986393)

### Establishment

Name	Address	ID/FEI	Business Operations
Northwind Pharmaceuticals, LLC		036986393	repack(51655-231)