## FEXOFENADINE HYDROCHLORIDE- fexofenadine hydrochloride tablet Pioneer Life Sciences, LLC Allergy Relief Fexofenadine Hydrochlroride Tablets 180 mg **Drug Facts** Active ingredient (in each caplet) Fexofenadine HCl 180 mg **Purpose Antihistamine** Uses temporarily relieves these symptoms due to hay fever or other upper respiratory $\sqcap$ runny nose $\sqcap$ sneezing $\sqcap$ itchy, watery eyes $\sqcap$ 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 $\square$ do not take more than directed $\square$ do not take at the same time as aluminum or magnesium antacids $\square$ 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

**Directions** 

Poison Control Center (1-800-222-1222) right away.

age uo not use

adults 65 years of age and older

consumers with kidney disease

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.

Questions or comments? Call toll-free 1 (732) 698-5070 Monday through Friday 9 am to 5 pm EST.

or visit: www.gencare.health.

This product is not manufactured or distributed by Chattem, Inc, a Sanofi Company, owner of the registered trademark Allegra® Allergy.

Distributed by: **Gencare Consumer Products, LLC** 40E Cotters Ln, Suite A, East Brunswick, NJ 08816

NDC 72090-010-01- 90 caplets



Size: 2" x 5" • Dec 04, 2023 • Peel Off Label

directed ■ do not take at the same time as aluminum or Glycol/macrogol, Povidone, Pregelatinized Starch, Red iron oxide, Silica, Titanium dioxide, Yellow iron oxide. Stop use and ask doctor if an allergic reaction to this CHILDREN. In case of overdose, get medical help or Children under 12 years of age. Do not use. Adults 65 years of age and older. Ask a doctor. Consumers with Inactive Ingredients: Colloidal Silicon Dioxide If pregnant or breast feeding, ask a health professional before use. KEEP OUT OF REACH OF and over. Take one 180 mg tablet with water once a Directions: Adults and children 12 years of age Croscarmellose Sodium Hypromellose, Magnesium magnesium antacids and not take with fruit juices contact a Poison Control Center (1-800-222-1222) Stearate, Microcrystalline Cellulose, Polyethylene day; do not take more than 1 tablet in 24 hours. product occurs. Seek medical help right away. ■ Store between 20 to 25°C (68°to 77°F)
■ Protect from excessive moisture

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Other Information:

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## FEXOFENADINE HYDROCHLORIDE

right away.

fexofenadine hydrochloride tablet

(see Directions)

When using this product: do not take more than

Facts (continued)

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72090-010
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII: E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	180 mg	

Inactive Ingredients		
Ingredient Name	Strength	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)		
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)		
POVIDONE (UNII: FZ989GH94E)		
STARCH, CORN (UNII: O8232NY3SJ)		
FERRIC OXIDE RED (UNII: 1K09F3G675)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

Product Characteristics			
Color	orange	Score	no score
Shape	CAPSULE	Size	17mm
Flavor		Imprint Code	
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72090-010- 01	90 in 1 BOTTLE; Type 0: Not a Combination Product	12/07/2023	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA210137	12/07/2023	

## Labeler - Pioneer Life Sciences, LLC (014092742)

Revised: 12/2023 Pioneer Life Sciences, LLC