ALLERGY RELIEF- fexofenadine hydrochloride tablet
AAA Pharmaceutical, Inc.

RES - 1192A - 2019-0911

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 right away (1-800-222-1222).

Directions

adults and children	take one 180 mg tablet with water once a

12 years of age and over	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 contains:** sodium 8.2 mg
- store between 20-25°C (68-77°F)
- protect from excessive moisture
- this product meets the requirements of USP *Dissolution Test 2*
- retain carton for complete product information and warnings

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, yellow iron oxide

Questions or Comments?

1-844-705-4384

PRINCIPAL DISPLAY PANEL

RESTORE U

NDC 57344-192-02

COMPARE TO THE ACTIVE INGREDIENT IN ALLEGRA® ALLERGY 24 HOUR

NON-DROWSY

Allergy Relief

Fexofenadine Hydrochloride Tablets, 180 mg / Antihistamine

Indoor and Outdoor Allergies – Original Prescription Strength

Relieves: Sneezing, Runny Nose, Itchy, Watery Eyes, Itchy Nose or Throat

24 HR

Actual Size

5 TABLETS

RESTORE U

NON-DROWSY

5 TABLETS

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Fexofenadine Hydrochloride Tablets, 180 mg

Antihistamine

NDC 57344-192-02

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NC

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Drug Facts (continued)

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Questions or comments? 1-844-705-4384

Distributed by:
AAA Pharmaceutical, Inc.
681 Main Street
Lumberton, NJ 08048
Made in India

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

DO NOT USE IF BLISTER UNITS ARE TORN OR BROKEN

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ALLERGY RELIEF

fexofenadine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:57344-192
Route of Administration	ORAL		

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

Inactive Ingredients	
Ingredient Name	**Strength**
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	

Product Characteristics			
Color	pink	**Score**	no score
Shape	OVAL (Modified oval shape)	**Size**	17mm
Flavor		**Imprint Code**	SG;202
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:57344-192-02	1 in 1 CARTON	10/01/2018	10/31/2022
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:57344-192-04	1 in 1 CARTON	06/01/2018	05/31/2021
2		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:57344-192-06	1 in 1 CARTON	06/01/2018	02/28/2023
		45 in 1 BOTTLE, PLASTIC; Type 0: Not a		

3		45 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204507	06/01/2018	02/28/2023

Labeler - AAA Pharmaceutical, Inc. (181192162)

Revised: 11/2021 AAA Pharmaceutical, Inc.