ALLERGY RELIEF- fexofenadine hydrochloride tablet AAA PHARMACEUTICAL, INC.

1192B-RES-2021-0706

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

	day; do not take more than 1 tablet in 24	
over	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

- store between 20-25°C (68-77°F)
- protect from excessive moisture
- retain carton for complete product information and warnings

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, lactose monohydrate, macrogol, magnesium stearate, microcrystalline cellulose, pregelatinized starch, titanium dioxide

Questions or comments?

1-844-705-4384

PRINCIPAL DISPLAY PANEL

Restore U

NDC 57344-292-04

†COMPARE TO THE ACTIVE INGREDIENT IN ALLEGRE® 24 HOUR

NON-DROWSY

Allergy Relief

Fexofenadine Hydrochloride Tablets, 180 mg / Antihistamine

Indoor / Outdoor Allergies

Relieves:

- Sneezing
- Runny Nose
- Itchy, Watery Eyes
- Itchy Nose or Throat

Actual Size

30 TABLETS



ALLERGY RELIEF						
fexofenadine hydrochloride ta	blet					
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:5734	NDC:57344-292	
Route of Administration	ORAL					
Active Ingredient/Active	Moiety					
Ingre	dient Name		Basis of Strength		Strength	
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582L0H6V)		FEXOFENADINE HYDROCHLORIDE		180 mg		
Inactive Ingredients						
_	Ingredient Name			St	rength	
CROSCARMELLOSE SODIUM (UN	-					
FERROSOFERRIC OXIDE (UNII: XM	10M87F357)					
FERRIC OXIDE RED (UNII: 1K09F3)	G675)					
FERRIC OXIDE YELLOW (UNII: EX	43802MRT)					
HYPROMELLOSE 2910 (15 MPA.	S) (UNII: 36SFW2JZ0W)					
LACTOSE MONOHYDRATE (UNII:	EWQ57Q8I5X)					
MAGNESIUM STEARATE (UNII: 70	097M6I30)					
MICROCRYSTALLINE CELLULOSI						
POLYETHYLENE GLYCOL 6000 (UNII: 30IQX730WE)					

SILICON DIOXI	DE (UNII: ETJ7Z6XBU4)						
STARCH, CORN	I (UNII: 08232NY3SJ)						
TITANIUM DIO	(IDE (UNII: 15FIX9V2JP)						
Product Characteristics							
Color	orange ((PEACH))	Score	no score				
Shape	OVAL (Capsule-shaped)	Size	17mm				
Flavor		Imprint Code	G6				

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57344- 292-03	1 in 1 CARTON	07/06/2021	
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:57344- 292-04	1 in 1 CARTON	07/06/2021	
2		30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:57344- 292-06	1 in 1 CARTON	07/06/2021	
3		45 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:57344- 292-02	1 in 1 CARTON	07/06/2021	
4		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
Μ	larketing	Information		
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	DA	ANDA211075	07/06/2021	

Labeler - AAA PHARMACEUTICAL, INC. (181192162)

Revised: 4/2022

AAA PHARMACEUTICAL, INC.