

**FEXOFENADINE HYDROCHLORIDE- fexofenadine hydrochloride tablet, film coated  
Bryant Ranch Prepack**

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**Perrigo Fexofenadine Hydrochloride Tablets, 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
- 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	take one 180 mg tablet with water once a day; do not take more than
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and over	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

- do not use if printed foil under cap is broken or missing
- store at 20°-25°C (68°-77°F)
- protect from excessive moisture
- this product meets the requirements of USP Dissolution Test 3

### Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, titanium dioxide

### Questions or comments?

1-800-719-9260

### HOW SUPPLIED

Product: 71335-0721

NDC: 71335-0721-1 30 TABLET, FILM COATED in a BOTTLE

NDC: 71335-0721-2 15 TABLET, FILM COATED in a BOTTLE

NDC: 71335-0721-3 60 TABLET, FILM COATED in a BOTTLE

NDC: 71335-0721-4 90 TABLET, FILM COATED in a BOTTLE

NDC: 71335-0721-5 5 TABLET, FILM COATED in a BOTTLE

NDC: 71335-0721-6 180 TABLET, FILM COATED in a BOTTLE

NDC: 71335-0721-7 100 TABLET, FILM COATED in a BOTTLE

### Fexofenadine Hcl 180mg Tablet

Packaged by Bryant Ranch

Burbank, CA 91504

# Fexofenadine Hcl 180mg Tablet

LOT  
117404

ORANGE (peach) ROUND 93; 7253

Store at room temp of  
20-25 C (68-77F)

Keep all drugs out of  
reach of children

Compare To:

Allegra 180mg Tablet

Perrigo

# 30

Exp: MM/YY

RX Only



NDC 7133507211

031901117404

## FEXOFENADINE HYDROCHLORIDE

fexofenadine hydrochloride tablet, film coated

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71335-0721(NDC:45802-571)
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FEXO FENADINE HYDRO CHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	180 mg

### Inactive Ingredients

Ingredient Name	Strength
SILICON DIO XIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERROSO FERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TITANIUM DIO XIDE (UNII: 15FIX9V2JP)	

### Product Characteristics

Color	ORANGE (peach)	Score	no score
Shape	ROUND	Size	12mm
Flavor		Imprint Code	93;7253

**Contains****Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71335-0721-3	60 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2012	
2	NDC:71335-0721-6	180 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2012	
3	NDC:71335-0721-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2012	
4	NDC:71335-0721-4	90 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2012	
5	NDC:71335-0721-7	100 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2012	
6	NDC:71335-0721-5	5 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2012	
7	NDC:71335-0721-2	15 in 1 BOTTLE; Type 0: Not a Combination Product	02/09/2012	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076447	04/22/2011	

**Labeler** - Bryant Ranch Prepack (171714327)**Establishment**

Name	Address	ID/FEI	Business Operations
Bryant Ranch Prepack		171714327	REPACK(71335-0721) , RELABEL(71335-0721)

Revised: 1/2020

Bryant Ranch Prepack