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

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 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

- do not use if carton is opened or printed foil under cap is broken or missing
- store between 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

NDC: 71335-0721-1: 30 Tablets in a BOTTLE

NDC: 71335-0721-2: 15 Tablets in a BOTTLE

NDC: 71335-0721-3: 60 Tablets in a BOTTLE

NDC: 71335-0721-4: 90 Tablets in a BOTTLE

NDC: 71335-0721-5: 5 Tablets in a BOTTLE

NDC: 71335-0721-6: 180 Tablets in a BOTTLE

NDC: 71335-0721-7: 100 Tablets in a BOTTLE

Fexofenadine Hcl 180mg Tablet

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**
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	180 mg

Inactive Ingredients	
Ingredient Name	**Strength**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERROSOFERRIC 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: 3WQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TITANIUM DIOXIDE (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:71335-0721-1	30 in 1 BOTTLE; Type 0: Not a Combination Product	03/09/2018	
2	NDC:71335-0721-2	15 in 1 BOTTLE; Type 0: Not a Combination Product	02/19/2019	
3	NDC:71335-0721-3	60 in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2021	
4	NDC:71335-0721-4	90 in 1 BOTTLE; Type 0: Not a Combination Product	02/05/2019	
5	NDC:71335-0721-5	5 in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2021	
6	NDC:71335-0721-6	180 in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2021	
7	NDC:71335-0721-7	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/27/2021	

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)

Registrant - Bryant Ranch Prepack (171714327)

Establishment

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