

FEXOFENADINE HYDROCHLORIDE- fexofenadine hydrochloride tablet, film coated
Carilion Materials Management

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

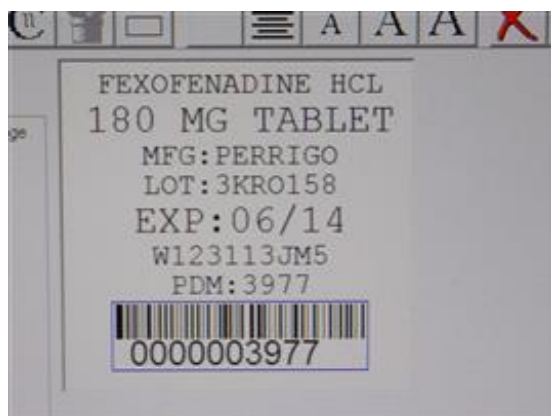
Inactive ingredients

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

Fexofenadine HCL 180 MG TAB



FEXOFENADINE HYDROCHLORIDE

fexofenadine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68151-3977(NDC:45802-571)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE -	FEXOFENADINE	180 mg

UNII:E6582LOH6V)			HYDROCHLORIDE		100 mg
Inactive Ingredients					
Ingredient Name				Strength	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)					
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)					
HYPROMELLOSES (UNII: 3NXW29V3WO)					
FERROSO FERRIC OXIDE (UNII: XM0M87F357)					
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)					
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)					
MAGNESIUM STEARATE (UNII: 70097M6I30)					
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)					
POLYETHYLENE GLYCOLS (UNII: 3WJQ0SDW1A)					
POVIDONES (UNII: FZ989GH94E)					
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)					
FERRIC OXIDE RED (UNII: 1K09F3G675)					
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:68151-3977-7	1 in 1 PACKAGE; Type 0: Not a Combination Product	04/22/2011		
Marketing Information					
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date	
ANDA	ANDA076447		04/22/2011		

Labeler - Carilion Materials Management (079239644)

Establishment

Name	Address	ID/FEI	Business Operations
Carilion Materials Management		079239644	REPACK(68151-3977)