

FEXOFENADINE HYDROCHLORIDE- fexofenadine hydrochloride tablet, film coated
Western Family Foods Inc

Western Family 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 blister unit is broken or torn (Use for Blister Configurations Only)
- do not use if printed foil under cap is broken or missing (Use for Bottle Configurations Only)
- 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

Package/Label Principal Display Panel

ORIGINAL PRESCRIPTION STRENGTH

ALLERGY

NON-DROWSY

Fexofenadine Hydrochloride TABLETS, 180 mg

ANTIHISTAMINE

24 HOUR

INDOOR & OUTDOOR ALLERGIES

RELIEF OF: SNEEZING; RUNNY NOSE

ITCHY, WATERY EYES; ITCHY NOSE or THROAT

Actual Size

30 TABLETS

180 mg EACH

ONE MONTH SUPPLY

COMPARE TO ALLEGRA® ALLERGY ACTIVE INGREDIENT



ORIGINAL PRESCRIPTION STRENGTH
ALLERGY

NON-DROWSY

Fexofenadine Hydrochloride

TABLETS, 180 mg
ANTIHISTAMINE



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TABLETS
180 mg EACH
ONE MONTH SUPPLY

COMPARE TO ALLEGRA® ALLERGY ACTIVE INGREDIENT*

12796-A-PER



0 15400 12796 7

LOT NO.

EXP.

: 57139 3M C1

ACTUAL SIZE



Fexofenadine Hydrochloride Tablets, 180 mg Carton Image 1



Drug Facts

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



NON-DROWSY
Fexofenadine
Hydrochloride
TABLETS, 180 mg
ANTIHISTAMINE

24 HOUR

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Made in Israel



Fexofenadine Hydrochloride Tablets, 180 mg Carton Image 2

FEXOFENADINE HYDROCHLORIDE

fexofenadine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55312-571
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 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:55312-571-39	1 in 1 CARTON		
1		30 in 1 BOTTLE		
2	NDC:55312-571-33	1 in 1 CARTON		
2		60 in 1 BOTTLE		
3	NDC:55312-571-22	15 in 1 CARTON		
3		1 in 1 BLISTER PACK		
4	NDC:55312-571-01	1 in 1 CARTON		
4		70 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076447	04/14/2011	

Labeler - Western Family Foods Inc (192166072)

Revised: 3/2014

Western Family Foods Inc