FEXOFENADINE HYDROCHLORIDE - fexofenadine hydrochloride tablet, film coated
Magno-Humphries Labs, Inc.

Active ingredient(in each tablet)

Fexofenadine HCl USP, 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 aluminium or magnesium antacids
- do not take with fruit juices (see directions)

Stop use and ask 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 (1-800-222-1222) right away.

Directions

adults and children 12 years of age and over	1 tablet with water 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

- store between 20°and 25°C (68°and 77°F)
- protect from excessive moisture and light

Inactive ingredients

anhydrous lactose, colloidal silicon dioxide, corn starch,croscarmellose sodium, hypromellose, lactose monohydrate, polyethylene glycol 400, pregelatinized corn starch, red iron oxide, steric acid, titanium dioxide, and yellow iron oxide,

Questions or comments?

call toll-free 1-800-935-6737

Principal Display Panel

***Compare to 24 Hour Allegra® Allergy active ingredient**

Fexofenadine Hydrochloride Tablets USP, 180 mg/antihistamine

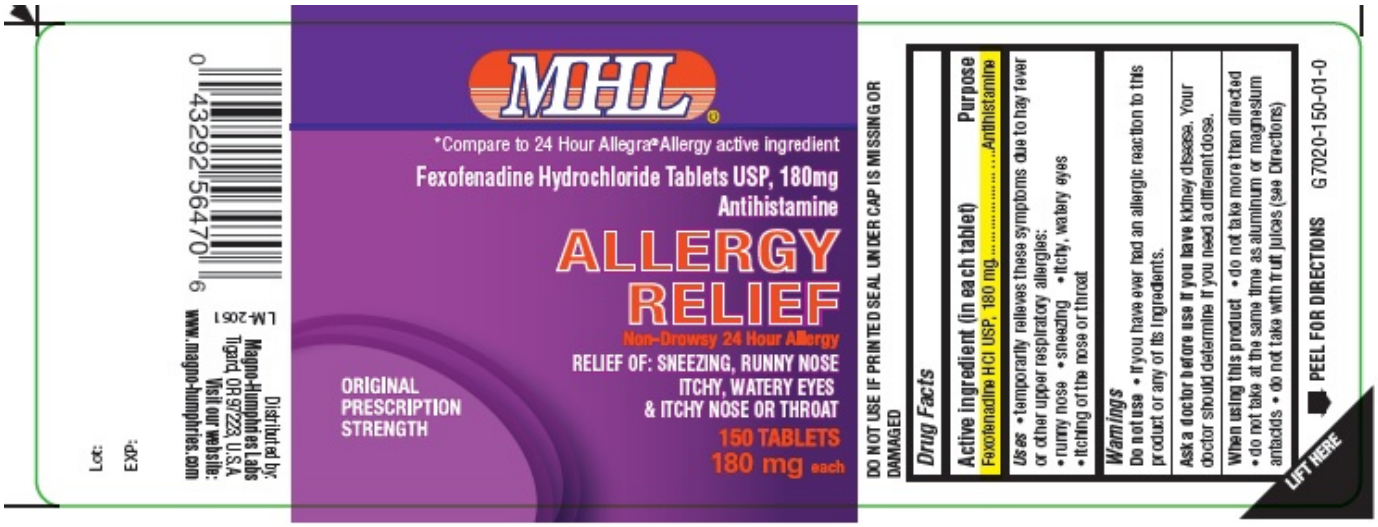
Allergy Relief
Non-Drowsy 24 hour Allergy

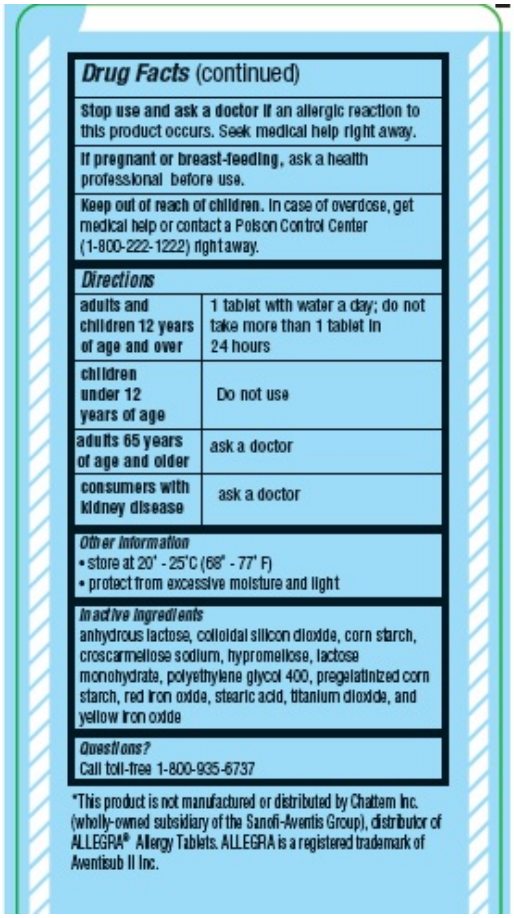
Relief of: Sneezing , Runny nose
Itchy, Watery Eyes &

Itchy Nose or Throat

Original Prescription Strength

150 Tablets 180 mg each





Drug Facts (continued)

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 (1-800-222-1222) right away.

Directions

adults and children 12 years of age and over	1 tablet with water 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
- store at 20° - 25°C (68° - 77° F)
- protect from excessive moisture and light

Inactive ingredients
anhydrous lactose, colloidal silicon dioxide, corn starch, croscarmellose sodium, hypromellose, lactose monohydrate, polyethylene glycol 400, pregelatinized corn starch, red iron oxide, stearic acid, titanium dioxide, and yellow iron oxide

Questions?
Call toll-free 1-800-935-6737

*This product is not manufactured or distributed by Chattem Inc. (wholly-owned subsidiary of the Sanofi-Aventis Group), distributor of ALLEGRA® Allergy Tablets. ALLEGRA is a registered trademark of Aventisub II Inc.

FEXOFENADINE HYDROCHLORIDE

fexofenadine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:54257-001
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
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	

STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	ORANGE (Peach)	**Score**	no score
Shape	CAPSULE (Bevel Edge, Biconvex)	**Size**	17mm
Flavor		**Imprint Code**	E;44
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54257-001-29	150 in 1 BOTTLE; Type 0: Not a Combination Product	01/15/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA202039	01/15/2015	

Labeler - Magno-Humphries Labs, Inc. (063251433)

Establishment

Name	Address	ID/FEI	Business Operations
Aurolife Pharma, LLC		829084461	MANUFACTURE(54257-001)

Revised: 9/2019 Magno-Humphries Labs, Inc.