

FEXOFENADINE HYDROCHLORIDE - fexofenadine hydrochloride tablet, film coated
Wal-Mart Stores, Inc

ACTIVE INGREDIENT(S)

Fexofenadine HCl USP, 180 mg

PURPOSE

Antihistamine

USE(S)

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 right away.

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

- safety sealed: do not use if carton is opened or if individual blister units are torn or opened
- store between 20⁰ and 25⁰C (68⁰ and 77⁰F)
- 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 OR COMMENTS?

call 1-888-287-1915

PRINCIPAL DISPLAY PANEL

NDC 49035-995-62

Non-Drowsy
Original Prescription Strength

Fexofenadine HCl Tablets USP,
180 mg/Antihistamine

Allergy Relief

Indoor and Outdoor Allergies
Relief of:

- Sneezing
- Runny nose
- Itchy, watery eyes
- Itchy nose or throat

180 mg EACH 15 TABLETS

6.75"

GLUE - NO COATING

1/16"

1/16"

1/8"

Drug Facts

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 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.

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

- safety sealed; do not use if carton is opened or if individual blister units are torn or opened
- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture and light

NO COATING
ALLEGRA

NO COATING
ALLEGRA

Drug Facts (continued)**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 or comments? call 1-888-267-1915

TAMPER EVIDENT: INDIVIDUAL BLISTER UNIT SEALED FOR YOUR PROTECTION. DO NOT USE IF INDIVIDUAL BLISTER UNIT IS OPEN OR TORN. IMPORTANT: Read the directions and warnings before use. Keep the carton. It contains important information.

LM-677

Lot
Exp.



180 mg EACH TABLETS



Original Prescription Strength
Fexofenadine HCl tablets USP,
180 mg/Antihistamine
Allergy Relief
 Indoor and Outdoor Allergies

- Itchy nose or throat
- Itchy, watery eyes
- Runny nose
- Sneezing



equate™

Manufactured by: Wal-Mart Stores, Inc. Item No. 48-27716
 Distributed by: Wal-Mart Stores, Inc. Item No. 48-27716
 The following is the name of the manufacturer, distributor, or registrant of the product: Wal-Mart Stores, Inc. Item No. 48-27716
 Satisfaction guaranteed - Or, we'll refund it or give you your money back. For questions or comments or to report an undesired reaction or side effect, please call 1-888-267-1915.



Original Prescription Strength
Allergy Relief
Fexofenadine HCl tablets USP,
180 mg/Antihistamine
 15 Tablets

TAMPER EVIDENT: INDIVIDUAL BLISTER UNIT SEALED FOR YOUR PROTECTION. DO NOT USE IF INDIVIDUAL BLISTER UNIT IS OPEN OR TORN.

equate™

7/8"

FEXOFENADINE HYDROCHLORIDE

fexofenadine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-995
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
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
POLYETHYLENE GLYCOL 400 (UNII: B697894SGQ)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	

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:49035-995-62	3 in 1 CARTON	09/30/2015	
1	NDC:49035-995-60	5 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA202039	09/30/2015	

Labeler - Wal-Mart Stores, Inc (051957769)

Establishment

Name	Address	ID/FEI	Business Operations
Aurolife Pharma, LLC		829084461	MANUFACTURE(49035-995)

Revised: 9/2019

Wal-Mart Stores, Inc