

FEXOFENADINE HYDROCHLORIDE - fexofenadine hydrochloride tablet, film coated
Walgreen CO

Active ingredient(in each tablet)

Fexofenadine HCl USP, 180 mg

Fexofenadine HCl USP, 60 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

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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

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 printed foil inner seal on bottle is torn or opened
- 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 1-855-274-4122

Principal Display Panel- 60 mg 24 ct

NDC 0363-0095-24

NON-DROWSY ORIGINAL PRESCRIPTION STRENGTH

Wal-Fex

12 HOUR ALLERGY

Fexofenadine Hydrochloride Tablets USP, 60 mg/antihistamine

12 Hour, TABLETS

INDOOR & OUTDOOR ALLERGIES

Relief of Sneezing; Runny nose; Itchy, Watery Eyes & Itchy Nose or Throat

24 Tablets 60 mg each

LEBG862V

Drug Facts

Active ingredient (in each tablet)

Fexofenadine HCl USP, 60 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 60 mg tablet with water every 12 hours; do not take more than 2 tablets 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

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 1-855-274-4122

LM-3277

Lot:
Exp:

NON-DROWSY
ORIGINAL
PRESCRIPTION STRENGTH

Wal-Fex®

12 HOUR ALLERGY

FEXOFENADINE HYDROCHLORIDE TABLETS USP, 60 mg / ANTIHISTAMINE

12 HOUR TABLETS

INDOOR & OUTDOOR ALLERGIES

- Relief of sneezing; runny nose; itchy, watery eyes & itchy nose or throat

24 TABLETS
60 mg EACH

Walgreens

Compare to Allegra® Allergy active ingredient^{††}

NDC 0363-0095-24



[†]Walgreens Pharmacist Survey
^{††}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.

DISTRIBUTED BY: WALGREEN CO.
200 WILMOT RD., DEERFIELD, IL 60015

Walgreens
100% SATISFACTION GUARANTEED
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ITEM 878919 W00000-0000-0



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ORG0818-F
REV1018



Principal Display Panel- 180 mg 15 ct

NDC 0363-0097-53

NON-DROWSY ORIGINAL PRESCRIPTION STRENGTH

Wal-Fex

Fexofenadine Hydrochloride Tablets USP, 180 mg/antihistamine

24 Hour, TABLETS

INDOOR & OUTDOOR ALLERGIES

Relief of Sneezing; Runny nose; Itchy, Watery Eyes & Itchy Nose or Throat

15 Tablets 180 mg each

ORIG0818-F
REV:10/18

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 1-855-274-4122

Walgreens

NON-DROWSY
ORIGINAL
PRESCRIPTION STRENGTH

Wal-Fex[®]

Compare to Allegra[®] Allergy active ingredient^{††}

NDC 0363-0097-53

24 HOUR ALLERGY

FEXOFENADINE HYDROCHLORIDE TABLETS USP, 180 mg / ANTIHISTAMINE



24 HOUR TABLETS

INDOOR & OUTDOOR ALLERGIES

- Relief of sneezing; runny nose; itchy, watery eyes & itchy nose or throat

15 TABLETS
180 mg EACH



ACTUAL SIZE

Lot: EXP:

LM-3278

[†]Walgreens Pharmacist Survey
^{††}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[®] Inc.

DISTRIBUTED BY: WALGREEN CO.
200 WILMOT RD., DEERFIELD, IL 60015

ITEM 878921 W00000-0000-0

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Drug Facts

Active ingredient (in each tablet)
Fexofenadine HCl USP, 180 mg
Antihistamine

Purpose
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 seal: 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

fexofenadine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0097
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)	
CROSCARMELOSE 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:0363-0097-55	1 in 1 CARTON	01/15/2015	
1		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:0363-0097-53	1 in 1 CARTON	01/15/2015	
2		15 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	01/15/2015	

FEXOFENADINE HYDROCHLORIDE

fexofenadine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-0095
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FEXO FENADINE HYDRO CHLORIDE (UNII: 2S068B75ZU) (FEXO FENADINE - UNII:E6582LOH6V)	FEXO FENADINE HYDROCHLORIDE	60 mg

Inactive Ingredients

Ingredient Name	Strength
ANHYDRO US LACTOSE (UNII: 3SY5LH9PMK)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STARCH, 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,Bioconvex)	Size	12mm
Flavor		Imprint Code	E;42
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0095-24	1 in 1 CARTON	10/05/2016	
1		24 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	10/05/2016	

Labeler - Walgreen CO (008965063)

Establishment

Name	Address	ID/FEI	Business Operations
Aurolife Pharma, LLC		829084461	MANUFACTURE(0363-0095, 0363-0097)

Revised: 9/2019

Walgreen CO