WAL-FEX- fexofenadine hydrochloride tablet
Walgreens Company

Fexofenadine Hydrochloride Tablets USP,180 mg

Active ingredient(s)

Fexofenadine Hydrochloride USP, 180 mg

Purpose

Antihistamine

Use(s)

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 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

(blister only) safety sealed: do not use if carton is opened or if individual blister units are torn or opened

(bottles only) safety sealed: do not use if carton is opened or if printed foil inner seal on bottle is torn or missing

Storage

- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture
- this product meets the requirements of USP Dissolution Test 2

Inactive ingredients

colloidal silicon dioxide, corn starch, croscarmellose sodium, FD&C Red no. 40, hypromellose, iron oxide black,magnesium stearate,mannitol, polyethylene glycol, powdered cellulose and titanium dioxide

Questions?

Call **1-888-375-3784**

Bottle Carton Label

Blister Carton Label: 15 count

Blister Carton Label


NON-DROWSY
ORIGINAL PRESCRIPTION STRENGTH
Wal-Fex®
24 HOUR ALLERGY
FEXOFENADINE HYDROCHLORIDE TABLETS USP, 180 mg / ANTIHISTAMINE
Walgreens
NON-DROWSY
ORIGINAL
PRESCRIPTION STRENGTH
Wal-Fex®
Compare to Allegra® Allergy active ingredient†
NDC 0363-0764-29
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
24 HOUR
ACTUAL SIZE
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
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.
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)
product form excessive moisture
this product meets the requirements of USP Dissolution Test 4
Inactive ingredients
colloidal silicon dioxide, corn starch, croscarmellose sodium, FD&C Red no. 40, hypromellose, iron oxide black, magnesium stearate, menthol, polydextrose, polyethylene glycol, powdered cellulose and titanium dioxide
Questions? Call 1-888-375-3784
DISTRIBUTED BY: WALGREEN CO.
200 WILMOT RD., DEERFIELD, IL 60015
Walgreens 100% SATISFACTION GUARANTEED
walgreens.com ©2018 Walgreen Co.
MADE IN INDIA
Walgreens Pharmacist Recommended
Walgreens Pharmacist Survey
†The product is not manufactured or distributed by Chattem, Inc. (part of the Sanofi Group), distributor of Allegra®
Allergy 24 Hour Tablets. Allegra® is a registered trademark of Aventis U.S.L.L.C.
AUTO-REORDER & SAVE
Free shipping + 20% off regular price
Learn more at Walgreens.com/AutoReorder
ORGXXXX-X
LOT
ITEM 44 1298 W 000 00-0 000 -0
TAMPER VD BRT: INDIVIDUAL BLISTER UNIT SEALED TO PROVIDE EVIDENCE OF TAMPERING. DO NOT USE IF BLISTER BOARD IS OPEN OR TORN.
IMPORTANT: Read the directions and warnings before use. Keep the carton. It contains important information.

WAL-FEX

fexofenadine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-0784(NDC:55111-784)
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
magnesium stearate (UNII: 70097M6I30)	
mannitol (UNII: 3OWL53L36A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
polyethylene glycol 400 (UNII: B697894SGQ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	PINK	**Score**	no score
Shape	OVAL	**Size**	7mm
Flavor		**Imprint Code**	194;R
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-0784-30	1 in 1 CARTON	04/13/2011	
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:0363-0784-43	2 in 1 CARTON	04/13/2011	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:0363-0784-40	1 in 1 CARTON	04/13/2011	
3		40 in 1 BOTTLE; Type 0: Not a Combination Product		

4	NDC:0363-0784-45	1 in 1 CARTON	04/13/2011	
4		45 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:0363-0784-75	1 in 1 CARTON	04/13/2011	
5		70 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:0363-0784-15	1 in 1 CARTON	04/13/2011	
6		150 in 1 BOTTLE; Type 0: Not a Combination Product		
7	NDC:0363-0784-90	1 in 1 CARTON	04/13/2011	
7		90 in 1 BOTTLE; Type 0: Not a Combination Product		
8	NDC:0363-0784-07	1 in 1 CARTON	04/13/2011	
8		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
9	NDC:0363-0784-29	3 in 1 CARTON	04/13/2011	
9		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
10	NDC:0363-0784-18	180 in 1 BOTTLE; Type 0: Not a Combination Product	07/13/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076502	04/13/2011	

Labeler - Walgreens Company (008965063)

Revised: 1/2022

Walgreens Company