FEXOFENADINE HYDROCHLORIDE- fexofenadine hydrochloride tablet
CVS Health Corp.

Fexofenadine HCl Tablets USP

Active ingredient (in each tablet)

Fexofenadine HCl USP, 180 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or otherupper 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

- (Bottle only)-safety sealed: do not use if carton is opened or if printed foil inner seal on bottle is torn or missing
- (Blister only)- Safety sealed: Do not use if seal is broken or if individual blister unit is torn or open
- 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, powder cellulose and titanium dioxide.

Questions?

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

Carton label

Carton Label: 40 count

CVS Health

Compare to the active ingedient in
Allegra® Allergy 24 Hour Tablets*

Indoor & Outdoor Alllergies

Original Prescription Strength

Allergy Relief

FEXOFENADINE HYDROCHLORIDE TABLETS USP, 180 MG

24 Hour Relief of:

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

Non-Drowsy 24 HOUR

Actual Bottle Size on Bottom Panel

Package Contains One Bottle

4**0 TABLETS 180 mg EACH**

CVSHealth

Indoor & Outdoor Allergies

Original Prescription Strength

Allergy Relief

FEXOFENADINE HYDROCHLORIDE
TABLETS USP, 180 mg
Antihistamine

33% MORE FREE 40 FOR THE PRICE OF 30

CVSHealth

Compare to the active ingredient in Allegra® Allergy 24 Hour Tablets*

Indoor & Outdoor Allergies

Original Prescription Strength

Allergy Relief

FEXOFENADINE HYDROCHLORIDE
TABLETS USP, 180 mg
Antihistamine

24 Hour Relief of:

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

24 HOUR

Non-Drowsy

Actual Bottle Size on Bottom Panel
Package Contains One Bottle

40 TABLETS 180 mg EACH

Actual Size

TAMPER EVIDENT: DO NOT USE IF FOIL SEAL UNDER CAP PRINTED WITH "SEALED FOR YOUR PROTECTION" IS BROKEN OR MISSING

IMPORTANT: Read the directions and warnings before use. Keep the carton. It contains important information.

Distributed by: CVS Pharmacy, Inc.
One CVS Drive, Woonsocket, RI 02895

CVS.com® 1-800-SHOP CVS

Made in India
V-13008

CVS Quality
Money Back Guarantee

*This 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 Aventisub LLC.

EXP LOT

#85200

Actual Size

Package Contains One Bottle

Drug Facts

Active ingredient (in each tablet)	Purpose
Fexofenadine HCl USP, 180 mg	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 printed foil inner seal on bottle is torn or missing
- store between 20° and 25°C (68° and 77°F)
- protect from 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, polyethylene glycol, powder cellulose and titanium dioxide

Questions? Call 1-888-375-3784

Bottle Label

Bottle Label: 40 count

CVS Health

Compare to the active ingedient in
Allegra® Allergy 24 Hour Tablets*

Indoor & Outdoor Alllergies

Original Prescription Strength

Allergy Relief

**FEXOFENADINE HYDROCHLORIDE
TABLETS USP, 180 MG**

24 Hour Relief of:

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

Non-Drowsy 24 HOUR

Actual Bottle Size on Bottom Panel

Package Contains One Bottle

4**0 TABLETS 180 mg EACH**

FEXOFENADINE HYDROCHLORIDE

fexofenadine hydrochloride tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69842-052(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:69842-052-30	1 in 1 CARTON	04/01/2011	
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:69842-052-90	1 in 1 CARTON	04/01/2011	
2		90 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:69842-052-40	1 in 1 CARTON	05/01/2013	
3		40 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:69842-052-45	1 in 1 CARTON	04/01/2011	
4		45 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:69842-052-60	2 in 1 CARTON	04/01/2011	
5		60 in 1 CARTON; Type 0: Not a Combination Product		

6	NDC:69842-052-18	1 in 1 CARTON	02/01/2013	
6		180 in 1 CARTON; Type 0: Not a Combination Product		
7	NDC:69842-052-02	1 in 1 CARTON	04/01/2015	
7		2 in 1 BLISTER PACK; Type 0: Not a Combination Product		
8	NDC:69842-052-07	1 in 1 CARTON	04/01/2011	
8		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
9	NDC:69842-052-15	3 in 1 CARTON	04/01/2011	
9		5 in 1 BLISTER PACK; Type 0: Not a Combination Product		
10	NDC:69842-052-29	3 in 1 CARTON	04/01/2011	
10		15 in 1 BLISTER PACK; Type 0: Not a Combination Product		
11	NDC:69842-052-70	1 in 1 CARTON	01/11/2019	
11		70 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076502	04/01/2011	

Labeler - CVS Health Corp. (062312574)

Revised: 12/2021 CVS Health Corp.