24HR ALLERGY RELIEF- fexofenadine hydrochloride tablet LEADER/ Cardinal Health 110, Inc.

Fexofenadine HCI Tablets USP

Active ingredient (in each tablet)

Fexofenadine HCI 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 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

	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, mannitol, polyethylene glycol, powder cellulose and titantium dioxide

Questions?

call **1-888-375-3784**

Package Label - 30 Count Carton

NDC 70000-0361-2 LEADER

NON-DROWSY

Original Prescription Strength

24HR Allergy Relief Fexofenadine Hydrochloride Tablets USP, 180 mg l Antihistamine

Indoor and Outdoor Allergies

24 Hour

Relief of

Sneezing, Runny Noise, Itchy Nose or Throat

and Itchy, Watery Eyes
COMPARE TO ALLEGRA®
ALLERGY 24HR
active ingredient*

100% Money Back Guarantee

30 TABLETS



NDC 70000-0361-2 LEADER

NON-DROWSY

Original Prescription Strength

24HR Allergy Relief Fexofenadine Hydrochloride Tablets USP, 180 mg | Antihistamine

Indoor and Outdoor Allergies

24 Hour

30 TABLETS



24HR ALLERGY RELIEF

fexofenadine hydrochloride tablet

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Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0361(NDC:55111-784)	
Route of Administration	ORAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
Fexofenadine Hydrochloride (UNII: 2S068B75ZU) (FEXOFENADINE -	Fexofenadine	100	

UNII:E6582LOH6V) Hydrochloride

Inactive Ingredients			
Ingredient Name	Strength		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)			
magnesium stearate (UNII: 70097M6I30)			
mannitol (UNII: 30WL53L36A)			
POWDERED CELLULOSE (UNII: SMD1X3X09M)			
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:70000- 0361-2	1 in 1 CARTON	04/27/2018		
1		30 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:70000- 0361-1	3 in 1 CARTON	04/27/2018		
2		5 in 1 BLISTER PACK; Type 0: Not a Combination Product			
3	NDC:70000- 0361-3	1 in 1 CARTON	04/27/2018		
3		70 in 1 BOTTLE; Type 0: Not a Combination Product			
4	NDC:70000- 0361-4	1 in 1 CARTON	04/27/2018		
4		90 in 1 BOTTLE; Type 0: Not a Combination Product			
5	NDC:70000- 0361-5	1 in 1 CARTON	04/23/2020		
5		180 in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information				
Marketing	Application Number or Monograph	Marketing Start	Marketing End	

Category	Citation	Date	Date
ANDA	ANDA076502	04/27/2018	

Labeler - LEADER/ Cardinal Health 110, Inc. (063997360)

Revised: 6/2021 LEADER/ Cardinal Health 110, Inc.