FEXOFENADINE HYDROCHLORIDE- fexofenadine hydrochloride tablet Strategic Sourcing Services, LLC

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

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, mannitol, polyethylene glycol, powder cellulose and titantium dioxide

Questions?

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

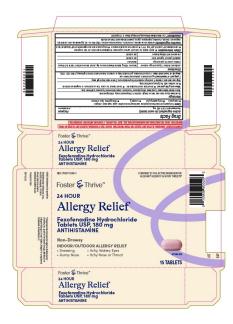
Carton Label

70ct Container and Container Carton labels





Carton



Container Label

FEXOFENADINE HYDROCHLORIDE

fexofenadine hydrochloride tablet

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70677-1009(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		
Strength		

polyethylene glycol 400 (UNII: B697894SGQ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
STARCH, CORN (UNII: 08232NY3SJ)	

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:70677- 1009-2	1 in 1 CARTON	05/05/2023		
1		30 in 1 BOTTLE; Type 0: Not a Combination Product			
2	NDC:70677- 1009-1	3 in 1 CARTON	05/05/2023		
2		5 in 1 BLISTER PACK; Type 0: Not a Combination Product			
3	NDC:70677- 1009-3	1 in 1 CARTON	05/05/2023		
3		70 in 1 BOTTLE; Type 0: Not a Combination Product			

Marketing Information			
Marketing Application Number or Monograph Category Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA076502	09/09/2019	

Labeler - Strategic Sourcing Services, LLC (116956644)

Revised: 4/2023 Strategic Sourcing Services, LLC