FEXOFENADINE HCL- fexofenadine hcl tablet Rite Aid
Non-drowsy
Fexofenadine HCI Tablets USP, 180mg
Antihistamine
Indoor/Outdoor Allergy relief
 sneezing runny nose itchy, watery eyes itchy nose or throat
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 take one 180 mg tablet with water once a day; do not take more than 1 tablet in 24 hours years of age and over

children under do not use 12 years of age

adults 65 years ask a doctor of age and older

consumers with ask a doctor kidney disease

Other information

] safety sealed: do not use if imprinted foil under bottle cap is opened	or	torn
store between 20° and 25°C (68° and 77°F)		
protect from excessive moisture		

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, lactose monohydrate, macrogol, magnesium stearate, microcrystalline cellulose, pregelatinized starch, titanium dioxide

Questions or comments?

contact 1-877-770-3183 Mon-Fri 8:00 AM EST to 5:00 PM PST

PDP



FEXOFENADINE HCL

fexofenadine hcl tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-7661
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
MAGNESIUM STEARATE (UNII: 70097M6I30)	
HYPROMELLOSE 2910 (15 MPA.S) (UNII: 36SFW2JZ0W)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL 6000 (UNII: 30IQX730WE)	
STARCH, CORN (UNII: O8232NY3SJ)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	

Product Characteristics			
Color	orange ((PEACH))	Score	no score
Shape	OVAL ((Capsule-shaped))	Size	17mm
Flavor		Imprint Code	G6
Contains			

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:11822- 7661-5	766 in 1 BOTTLE; Type 0: Not a Combination Product	10/25/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211075	10/25/2024	

Labeler - Rite Aid (014578892)

Revised: 12/2024 Rite Aid