# FEXOFENADINE HCL- fexofenadine hcl tablet TARGET CORPORATION

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### Fexofenadine HCL USP 180mg

#### **Antihistamine**

### 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 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

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 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?**

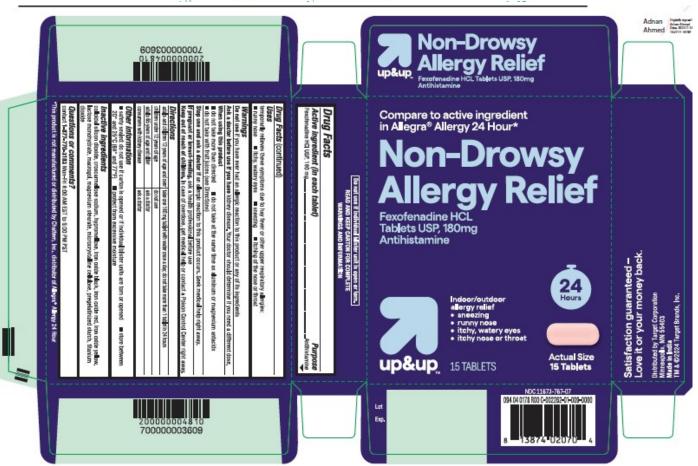
contact **1-877-770-3183** Mon-Fri 8:00 AM EST to 5:00 PM PST

#### **PDP**









### **FEXOFENADINE HCL**

fexofenadine hcl tablet

### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11673-767

Active Ingredient/Active Moiety			
Ingredient Name	<b>Basis of Strength</b>	Strength	
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII: E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	180 mg	

Inactive Ingredients		
Ingredient Name	Strength	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
FERRIC OXIDE YELLOW (UNII: EX43802MRT)		
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)		
FERRIC OXIDE RED (UNII: 1K09F3G675)		
HYPROMELLOSE 2910 (15 MPA.S) (UNII: 36SFW2JZ0W)		
STARCH, CORN (UNII: O8232NY3SJ)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
POLYETHYLENE GLYCOL 6000 (UNII: 30IQX730WE)		

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

ı	Packaging				
4	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:11673- 767-07	15 in 1 BLISTER PACK; Type 0: Not a Combination Product	03/01/2024		
:	NDC:11673- 767-03	30 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2024		
3	NDC:11673- 767-34	70 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2024		
4	NDC:11673- 767-15	150 in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2024		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211075	03/01/2024	

## Labeler - TARGET CORPORATION (006961700)

Revised: 12/2023 TARGET CORPORATION