ALLERGY RELIEF- fexofenadine hcl tablet, coated TARGET Corporation

DRUG FACTS

Active ingredient (in each film-coated 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 years of age and over	take one 18 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

- each tablet contains: sodium 8 mg
- store between 20-25°C (68-77°F)
- protect from excessive moisture
- this product meets the requirements of USP Dissolution Test 2

Inactive ingredients

anhydrous lactose, colloidal silicon dioxide, corn starch, croscarmellose sodium, lactose monohydrate, pregelatinized starch (maize), stearic acid, opadry pink 03B84893 containing hypromellose, polyethylene glycol, red iron oxide, titanium dioxide, and yellow iron oxide

Questions or comments?

call **1-800-910-6874**

Principal Display Panel

Compare to the active ingredient in Allegra® Allergy 24 hour*

non-drowsy

Allergy Relief

fexofenadine HCI 180mg / antihistamine

original prescription strength

indoor and outdoor allergies

Relief of:

- sneezing
- runny nose
- itchy, watery eyes
- itchy nose or throat

COATED CAPLETS** (**Capsule-shaped tablets)

*This product is not manufactured or distributed by Chattem Inc., distributor of Allegra® Allergy 24 Hour.

TAMPER EVIDENT: DO NOT USE IF CARTON IS OPENED OR IF BLISTER UNIT IS TORN, BROKEN OR SHOWS ANY SIGNS OF TAMPERING.

KEEP OUTER CARTON FOR COMPLETE WARNINGS AND PRODUCT INFORMATION.

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Product Label

Compare to active ingredient in Allegra® Allergy 24 Hour* NDC 11673-104-30 non-drowsy /antihistamine fexofenadine HCI original prescription strength indoor and outdoor allergies relief of: sneezing
itchy, watery eyes
runny nose
itchy throat or nose upaup



CAPLETS**

30 + 10 = 40 COATED CAPLETS** (**Capsule-shaped tablets)



bonus

10 more!



non-drowsy

fexofenadine HCI /antihistamine





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NOTES OUTER CARTON FOR COMPLETE WANNIES AND PRODUCT INFORMATION.

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(in each film-coated tablet)

TARGET Non-Drowsy Allergy Relief

ALLERGY RELIEF

fexofenadine hcl tablet, coated

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11673-104

Route of Administration ORAL

Active Ingredient/Active Moiety

Ingredient Na	ame	Basis of Strength	Strength
FEXOFENADINE HYDROCHLORIDE (UNII: 25	5068B75ZU) (FEXOFENADINE -	FEXOFENADINE	180 ma

UNII:E6582LOH6V) HYDROCHLORIDE 160 HIG

Inactive Ingredients			
Ingredient Name	Strength		
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)			
HYPROMELLOSES (UNII: 3NXW29V3WO)			
FERRIC OXIDE RED (UNII: 1K09F3G675)			
FERRIC OXIDE YELLOW (UNII: EX43802MRT)			
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)			
STARCH, CORN (UNII: O8232NY3SJ)			
STEARIC ACID (UNII: 4ELV7Z65AP)			

Product Characteristics			
Color	white	Score	no score
Shape	CAPSULE	Size	17mm
Flavor		Imprint Code	SG;202
Contains			

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673- 104-15	1 in 1 CARTON	01/11/2018	05/30/2025
1		15 in 1 BLISTER PACK; Type 0: Not a Combination Product		
_	NDC:11673-	1 :- 1 DOV	01/11/2010	05/20/2025

4	104-12	I III I DOV	NT/TT/TNTO	U3/3U/ZUZ3
2		120 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
3	NDC:11673- 104-70	1 in 1 BOX	01/11/2018	05/30/2025
3		70 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		
4	NDC:11673- 104-30	1 in 1 BOX	01/11/2018	05/30/2025
4		40 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA079112	01/11/2018	05/30/2025

Labeler - TARGET Corporation (006961700)

Revised: 11/2023 TARGET Corporation