

ALLERGY RELIEF- fexofenadine hcl tablet, coated
TARGET Corporation

DRUG FACTS

Active ingredient (in each film-coated 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 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 HCl 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.

Distributed by Target Corporation

Minneapolis, MN 55403

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Product Label

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 Name	Basis of Strength	Strength
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V)	FEXOFENADINE HYDROCHLORIDE	180 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
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		
1	NDC:11673-	1 in 1 BOX	01/11/2018	05/30/2025

2	104-12	1 in 1 BOX	01/11/2018	05/30/2025
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