UP AND UP ALLERGY RELIEF- fexofenadine hydrochloride tablet, film coated
Target Corporation

Target Corporation Allergy Relief Drug Facts

Active ingredient (in each tablet)

Fexofenadine HCl 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. (1-800-222-1222)

Directions

adults and children 12 years of age	take one 180 mg tablet with water once a day; do not take more than

and over	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

- do not use if blister unit is broken or torn (Use for Blister Configuration Only)
- do not use if printed foil under cap is broken or missing (Use for Bottle Configuration Only)
- store at 20°-25°C (68°-77°F)
- protect from excessive moisture
- this product meets the requirements of USP Dissolution Test 3

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, titanium dioxide

Questions?

Call 1-800-910-6874

Package/Label Principal Display Panel

original prescription strength

non-drowsy

allergy relief

fexofenadine hydrochloride tablets, 180 mg

antihistamine

Compare to active ingredient in Allegra® Allergy

relief of: sneezing/runny nose/itchy, watery eyes/itchy nose or throat

indoor & outdoor allergies

ACTUAL SIZE

24 HOUR

TABLETS {replace "#" with actual number of tablets per package}

180 mg EACH

original
prescription strength
non-drowsy
allergy relief
fexofenadine hydrochloride
tablets, **180 mg**/antihistamine

up&up™

NDC 11673-571-75

original prescription strength non-drowsy **allergy relief**

fexofenadine hydrochloride tablets, **180 mg**
antihistamine

Compare to active ingredient in Allegra® Allergy*

relief of: sneezing/runny nose/itchy, watery eyes/
itchy nose or throat

indoor & outdoor allergies

90
TABLETS

up&up™

90 TABLETS 180 mg EACH

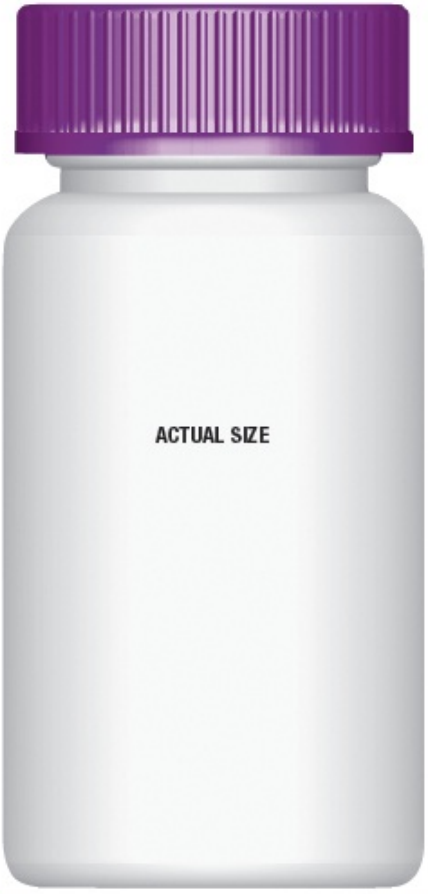
094 04 0087 ID132461
Distributed by Target Corporation
Minneapolis, MN 55403
Made in Israel

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

LOT NO.

EXP.

: 57175 UW C1



Drug Facts

Active Ingredient (in each tablet) — **Purpose**

Fexofenadine HCl 180 mg ..... Antihistamine

Uses

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- sneezing
- itching of the nose or throat

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adults 65 years of age and older	ask a doctor
consumers with kidney disease	ask a doctor

Other Information

- do not use if printed foil under cap is broken or missing
- store at 20°-25°C (68°-77°F)
- protect from excessive moisture
- this product meets the requirements of USP *Dissolution Test 3*

Inactive Ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, titanium dioxide

Questions? **Call 1-800-910-6874**

original
prescription
strength
non-drowsy

allergy relief

fexofenadine hydrochloride
tablets, **180 mg**/antihistamine

up&up™

UP AND UP ALLERGY RELIEF

fexofenadine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG LABEL	**Item Code (Source)**	NDC:11673-571
Route of Administration	ORAL	**DEA Schedule**	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FEXOFENADINE HYDROCHLORIDE (FEXOFENADINE)	FEXOFENADINE HYDROCHLORIDE	180 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE	
CROSCARMELLOSE SODIUM	
HYPROMELLOSES	
FERROSOFERRIC OXIDE	
FERRIC OXIDE YELLOW	
LACTOSE MONOHYDRATE	
MAGNESIUM STEARATE	
CELLULOSE, MICROCRYSTALLINE	
POLYETHYLENE GLYCOLS	
POVIDONES	
TITANIUM DIOXIDE	
FERRIC OXIDE RED	

Product Characteristics

Color	ORANGE (peach)	**Score**	no score
Shape	ROUND	**Size**	12mm
Flavor		**Imprint Code**	93;7253
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-571-39	1 in 1 CARTON		
1		30 in 1 BOTTLE		
2	NDC:11673-571-95	1 in 1 CARTON		
2		45 in 1 BOTTLE		
3	NDC:11673-571-22	3 in 1 CARTON		
3		5 in 1 BLISTER PACK		
4	NDC:11673-571-75	1 in 1 CARTON		
4		90 in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076447	04/13/2011	

Labeler - Target Corporation (006961700)

Revised: 4/2013 Target Corporation