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 and over	take one 180 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

- do not use if printed blister unit is broken or torn
- store between 20° -25°C (68° -77°F)
- protect from excessive moisture
- this product meets the requirements of USP *Dissolution Test 2*

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, FD&C blue #2 aluminum lake, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, pregelatinized starch, talc, titanium dioxide

Questions?

Call 1-888-547-7400

Package/Label Principal Display Panel

Compare to active ingredient in Allegra® Allergy

non-drowsy

allergy relief

fexofenadine hydrochloride tablets 180 mg/antihistamine

indoor/outdoor allergy relief

• sneezing

• runny nose

• itchy, watery eyes

• itchy nose or throat

ACTUAL SIZE

24 HOUR

15 TABLETS

15 TABLETS

CONVENIENT RECLOSING TAB

OPEN OTHER END

84722 UW C1

GLUTEN FREE

Drug Facts

Active ingredient (in each tablet)	Purpose
Fexofenadine HCl 180 mg	Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
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Warnings

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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 (1-800-22-1222)

Drug Facts (continued)

Directions

adults and children 12 years of age and over	take one 180 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

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Questions? Call 1-888-547-7400

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

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up&up™

non-drowsy

allergy relief

fexofenadine hydrochloride tablets **180 mg**/antihistamine

NDC 11673-617-22

Compare to active ingredient in Allegra® Allergy*

non-drowsy

allergy relief

fexofenadine hydrochloride tablets
180 mg/antihistamine

ACTUAL SIZE

up&up™

indoor/outdoor allergy relief

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

24 HOUR

15 TABLETS

15 TABLETS

UP AND UP ALLERGY RELIEF

fexofenadine hydrochloride tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-617
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**
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	PINK	**Score**	no score
Shape	OVAL	**Size**	18mm
Flavor		**Imprint Code**	L847
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:11673-617-22	15 in 1 CARTON	09/30/2021	04/30/2025
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:11673-617-39	1 in 1 CARTON	09/17/2021	12/31/2024
2		30 in 1 BOTTLE; Type 0: Not a Combination		

2		Product		
3	NDC:11673-617-00	1 in 1 CARTON	09/30/2021	09/30/2021
3		40 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:11673-617-01	1 in 1 CARTON	09/30/2021	06/30/2025
4		70 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:11673-617-47	1 in 1 CARTON	09/30/2021	08/31/2025
5		150 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA212971	09/17/2021	08/31/2025

Labeler - Target Corporation (006961700)

Revised: 6/2025 Target Corporation