TOPCARE ALLERGY RELIEF- fexofenadine hydrochloride tablet, film coated
Topco Associates LLC

Topco Associates LLC. 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 or comments?

1-888-423-0139

Package/Label Principal Display Panel

COMPARE TO ALLEGRA® ALLERGY ACTIVE INGREDIENT

NON-DROWSY

Allergy Relief

FEXOFENADINE HYDROCHLORIDE TABLETS, 180 mg (ANTIHISTAMINE)

24 HR

INDOOR/OUTDOOR ALLERGY RELIEF

- Sneezing – Runny Nose

- Itchy, Watery Eyes

- Itchy Nose or Throat

5 TABLETS

actual size

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Drug Facts (continued)

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Questions or comments? 1-888-423-0139

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topcare@topco.com www.topcarebrand.com

GF GLUTEN FREE

847 13 88 C1

QUALITY GUARANTEED

This TopCare® product is laboratory tested to guarantee its highest quality. Your total satisfaction is guaranteed.

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

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+TopCare® health

Allergy Relief

FEXOFENADINE HYDROCHLORIDE TABLETS, 180 mg (ANTIHISTAMINE)

Scan here for more information or call 1-888-423-0139

NDC 36800-691-13

+TopCare® health

COMPARE TO ALLEGRA® ALLERGY ACTIVE INGREDIENT*

NON-DROWSY

Allergy Relief

FEXOFENADINE HYDROCHLORIDE TABLETS, 180 mg (ANTIHISTAMINE)

24 HR

INDOOR/OUTDOOR ALLERGY RELIEF
- Sneezing • Runny Nose
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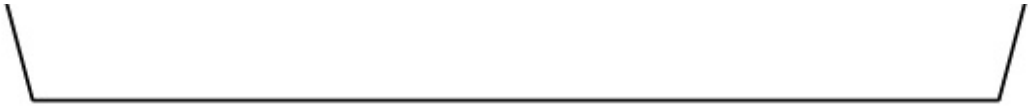

TOPCARE ALLERGY RELIEF

fexofenadine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:36800-691
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:36800-691-13	5 in 1 CARTON	08/27/2021	10/31/2023
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:36800-691-	1 in 1 CARTON	08/27/2021	

2	39	1 in 1 CARTON	08/23/2021	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:36800-691-22	15 in 1 CARTON	09/21/2021	09/21/2021
3		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
4	NDC:36800-691-49	1 in 1 CARTON	09/21/2021	
4		40 in 1 BOTTLE; Type 0: Not a Combination Product		
5	NDC:36800-691-95	1 in 1 CARTON	09/21/2021	12/31/2023
5		45 in 1 BOTTLE; Type 0: Not a Combination Product		
6	NDC:36800-691-75	1 in 1 CARTON	09/21/2021	
6		90 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	08/23/2021	

Labeler - Topco Associates LLC (006935977)

Revised: 12/2025 Topco Associates LLC