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



39800 49559



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Que stions or comments? 1-88-423 0139

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Active ingredient (in each tablet) Purpose Feorenatine HCl 180 mg.......Antihatamine

Storing Facts

8471388



Allergy Relief

FEXOFENADINE HYDROCHLORIDE TABLETS, 180 mg (ANTIHISTAMINE)



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

NDC36800-691-13

COMPARE TO ALLEGRA® ALLERGY ACTIVE INGREDIENT*



NON-DROWSY

Allergy Relief

FEXOFENADINE HYDROCHLORIDE TABLETS, **180 mg** (ANTIHISTAMINE)



INDOOR/OUTDOOR ALLERGY RELIEF

- Sneezing Runny Nose
- Itchy, Watery Eyes
- Itchy Nose or Throat

5 TABLETS

actual size

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

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Ingredient Name	Basis of Strength	Strength
	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: 3WJQ0SDW1A)			
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			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:36800-691- 13	5 in 1 CARTON	08/27/2021	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
_	NDC:36800-691-	1 to 1 CARTON	00/22/2021	

	39	I III I CARTON	00/23/2021	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:36800-691- 22	15 in 1 CARTON	09/21/2021	
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	
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: 9/2021 Topco Associates LLC