

**ALLERGY RELIEF- fexofenadine hydrochloride tablet, film coated
H E B**

HEB 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?

1-800-719-9260

Package/Label Principal Display Panel

Compare to Allegra[®] Allergy active ingredient

H-E-B[®]

Allergy Relief

Fexofenadine Hydrochloride Tablets, 180 mg/Antihistamine

NON-DROWSY

INDOOR/OUTDOOR ALLERGY RELIEF

24 HR

- Sneezing
- Runny Nose
- Itchy, Watery Eyes

- Itchy Nose or Throat

actual size

15 TABLETS



Allergy Relief

Fexofenadine Hydrochloride Tablets, 180 mg
Antihistamine



Compare to Allegra® Allergy active ingredient*

NDC 37808-847-22



Allergy Relief

Fexofenadine Hydrochloride
Tablets, 180mg/Antihistamine

NON-DROWSY

INDOOR/OUTDOOR ALLERGY RELIEF

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



actual size

15 TABLETS



84722 1101

OPEN OTHER END

Drug Facts

Active ingredient (in each tablet) Purpose
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 ■ sneezing ■ itching of the nose or throat

Warnings
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Drug Facts (continued)

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Questions or comments? 1-800-719-9260

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

GLUTEN FREE

CONVENIENT RECLOSING TAB



MADE WITH PRIDE & CARE FOR H-E-B® SAN ANTONIO, TX 78204

100% GUARANTEE promise

If you aren't completely pleased with this product, we'll happily replace it or refund your money. You have our word on it.

ALLERGY RELIEF

fexofenadine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37808-847
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: 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			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-847-22	15 in 1 CARTON	12/22/2021	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:37808-847-39	1 in 1 CARTON	12/22/2021	
3		30 in 1 BOTTLE; Type 0: Not a Combination		

4		Product		
3	NDC:37808-847-49	1 in 1 CARTON	12/30/2021	
3		40 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:37808-847-95	1 in 1 CARTON	12/22/2021	
4		45 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	12/22/2021	

Labeler - H E B (007924756)

Revised: 12/2021

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