ALLERGY RELIEF- fexofenadine hcl tablet, film coated H E B

HEB Allergy Relief Drug Facts

Active ingredient (in each tablet)

Fexofenadine HCI 60 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	take one 60 mg tablet with water every 12 hours; do
age and over	not take more than 2 tablets 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
- 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 or comments?

1-800-719-9260

Package/Label Principal Display Panel

Compare to Allegra® Allergy active ingredient

Allergy Relief

Fexofenadine Hydrochloride Tablets, 60 mg/Antihistamine

NON-DROWSY

INDOOR/OUTDOOR ALLERGY RELIEF

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

actual size

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help or contact a Poison Control Center right away

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product or any of its ingredients.

When using this product

■ do not take with fruit juices (see Directions)

should determine if you need a different dose.

Keep out of reach of children, in case of overdose, get medical If pregnantor breast-fee ding, ask a health professional before use.

Stop use and ask a doctoritan allergic reaction to this product

do not take at the same tim e as aluminum or magnesium an tacids

As k a doctor before use if you have kidney disease. Your doctor



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

UNGSTIONS OF COMMENTS? 1-800-719-9260

polye thyle ne glycol, povidone, titanium dioxide monolnyd rate, mag ne siu m stea rat e, microcry stall ine cellulose, iuou oxide black, iron oxide red, iro n oxide yello w, lactose colloidal silic on dioxi de , cro scarmel lose so dium , hyp ro mello se, STATE INGREDIES

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suq o ver 1 2 Je sız of siğe suq o ver	take one 60 mg table t with water every 12 hours; do not take more than 2 tablets in 24 hours
Directions	

■ ECHING OT The nose or throat ■ гил пу пове таку, watery е ува

up per respiratory all ergies: temporarily relieves these symptoms due to hay fever orother

Do not use if you have ever had an allergic reaction to this

Pexofe nadine HCl 60 mg..... ənimistənində.. Active ingredient (in each tablet) 9sod 1nd

Drug Facts

NDC 37808-425-53

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Drug Facts (continued)



Allergy Relief

Fexofenadine Hydrochloride Tablets, 60 mg/Antihistamine

Compare to Allegra® Allergy active ingredient*



Allergy Relief

Fexofenadine Hydrochloride Tablets, 60 mg/Antihistamine

NON-DROWSY

INDOOR/OUTDOOR ALLERGY RELIEF

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



actual

12 TABLETS

















ALLERGY RELIEF

fexofenadine hcl tablet, film coated

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:37808-425

Route of Administration ORAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - HYDROCHLORIDE UNII: E6582LOH6V) FEXOFENADINE HYDROCHLORIDE 60 mg

Inactive Ingredients	
Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	ORANGE (Peach)	Score	no score
Shape	ROUND	Size	8mm
Flavor		Imprint Code	93;7252
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-425- 53	12 in 1 CARTON	08/24/2011	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA076447	08/24/2011	

Labeler - H E B (007924756)

Revised: 7/2021 H E B