ALLERGY RELIEF- fexofenadine hydrochloride tablet Magno-Humphries, Inc.

MAGNO-HUMPHRIES - Antihistamine Allergy Relief

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
- sneezing
- itchy, watery eyes
- 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 (1-800-222-1222) right away.

Directions

adults and children 12 years of age and over children under 12 years of age

adults 65 years of age and older

consumers with kidney disease

take one 180 mg tablet with water once a day; do not take more than 1 tablet in 24 hours

do not use

ask a doctor

ask a doctor

Other information

- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture
- each tablet contains: sodium 8.2 mg
- this product meets the requirements of USP Dissolution Test 2

Inactive ingredients

anhydrous lactose, colloidal silicon dioxide, corn starch, croscarmellose sodium, hypromellose, lactose monohydrate, polyethylene glycol, pregelatinized starch (maize), red iron oxide, stearic acid, titanium dioxide, yellow iron oxide.

Questions?

call toll-free 1-800-935-6737



DO NOT USE IF PRINTED SEAL UNDER CAP IS MISSING OR DAMAGED

Drug Facts

Uses

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- runny nose
- sneezing
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- itching of the nose or throat

Warnings

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- 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 (1-800-222-1222) right away.

PEEL FOR DIRECTIONS

G7022-100-01-0

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 and older	ask a doctor
consumers with kidney disease	ask a doctor

Other information

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Questions? call toll-free 1-800-935-6737

*This product is not manufactured or distributed by the owner of the registered trademark Allegra® Allergy 24 Hour Tablets

> Distributed by: Magno-Humphries Labs Tigard, OR 97223 U.S.A.

Visit our website: www.magno-humphries.com

ALLERGY RELIEF

fexofenadine hydrochloride tablet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:54257-170

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
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
CORN (UNII: 0N86727070)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	

Product Characteristics

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Color	pink	Score	no score
Shape	CAPSULE	Size	18mm
Flavor		Imprint Code	SG202
Contains			

Packaging

- 1		gg			
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:54257-170- 02	100 in 1 BOTTLE; Type 0: Not a Combination Product	03/25/2021	

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
ANDA	ANDA204507	03/25/2021	

Revised: 1/2024 Magno-Humphries, Inc.