

ALLERGY RELIEF- fexofenadine hydrochloride tablet
Chain Drug Marketing Association, Inc.

1191A-QCH-2023-0522

Drug Facts

Active ingredient (in each tablet)

Fexofenadine HCl 60 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 right away (1-800-222-1222).

Directions

adults and children	take one 60 mg tablet with water every 12
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12 years of age and over	hours; do 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

- store between 20-25°C (68-77°F)
- protect from excessive moisture
- retain carton for complete product information and warnings

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, lactose monohydrate, macrogol, magnesium stearate, microcrystalline cellulose, pregelatinized starch, titanium dioxide

PRINCIPAL DISPLAY PANEL

QUALITY CHOICE®

NDC 83324-009-12

*Compare to the Active Ingredient in ALLEGRA® Allergy 12 Hour

Non-Drowsy

Allergy Relief

Fexofenadine Hydrochloride Tablets, 60 mg | Antihistamine

Indoor / Outdoor Allergies

Relief of:

Sneezing | Runny Nose

Itchy, Watery Eyes

Itchy Nose or Throat

12 Hour

actual size

12 TABLETS



Non-Drowsy

Allergy Relief



Fexofenadine Hydrochloride Tablets, 60 mg | Antihistamine

*This product is not manufactured or distributed by Chiltern, Inc., distributor of Allegra® Allergy 12 Hour.

DO NOT USE IF BLISTER UNITS ARE TORN OR BROKEN



Distributed by CDMA, Inc.
Novi, MI 48375
www.qualitychoice.com
Questions: 800-935-2362
Made in India

Drug Facts	
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Purpose 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	
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Non-Drowsy

Allergy Relief



Fexofenadine Hydrochloride Tablets, 60 mg | Antihistamine

NDC 83324-009-12



*Compare to the Active Ingredient in ALLEGRA® Allergy 12 Hour

Non-Drowsy

Allergy Relief

Fexofenadine Hydrochloride Tablets, 60 mg | Antihistamine

Indoor / Outdoor Allergies

Relief of:

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



actual size



12 TABLETS

NC

NC

NC

ALLERGY RELIEF

fexofenadine hydrochloride tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83324-009
Route of Administration	ORAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
FERROSFERRIC OXIDE (UNII: XM0M87F357)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX438O2MRT)	
HYPROMELLOSE 2910 (15 MPA.S) (UNII: 36SFW2JZ0W)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 6000 (UNII: 30IQX730WE)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	orange ((PEACH))	Score	no score
Shape	OVAL	Size	12mm
Flavor		Imprint Code	G5
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83324-009-12	1 in 1 CARTON	05/22/2023	
1		12 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:83324-009-24	2 in 1 CARTON	05/22/2023	
2		12 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	ANDA211075	05/22/2023	

Labeler - Chain Drug Marketing Association, Inc. (011920774)

Revised: 5/2023

Chain Drug Marketing Association, Inc.