

ALLERGY RELIEF- fexofenadine hydrochloride 60 mg tablet
Chain Drug Marketing Association INC

Fexofenadine HCl Tablets USP

Active ingredient(s)

Fexofenadine HCl USP, 60 mg

Purpose

Antihistamine

Use(s)

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 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 60 mg tablet with water every 12 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

- safety sealed: do not use if carton is opened or if printed foil inner seal on bottle is torn or missing
- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture
- this product meets the requirements of USP Dissolution Test 4

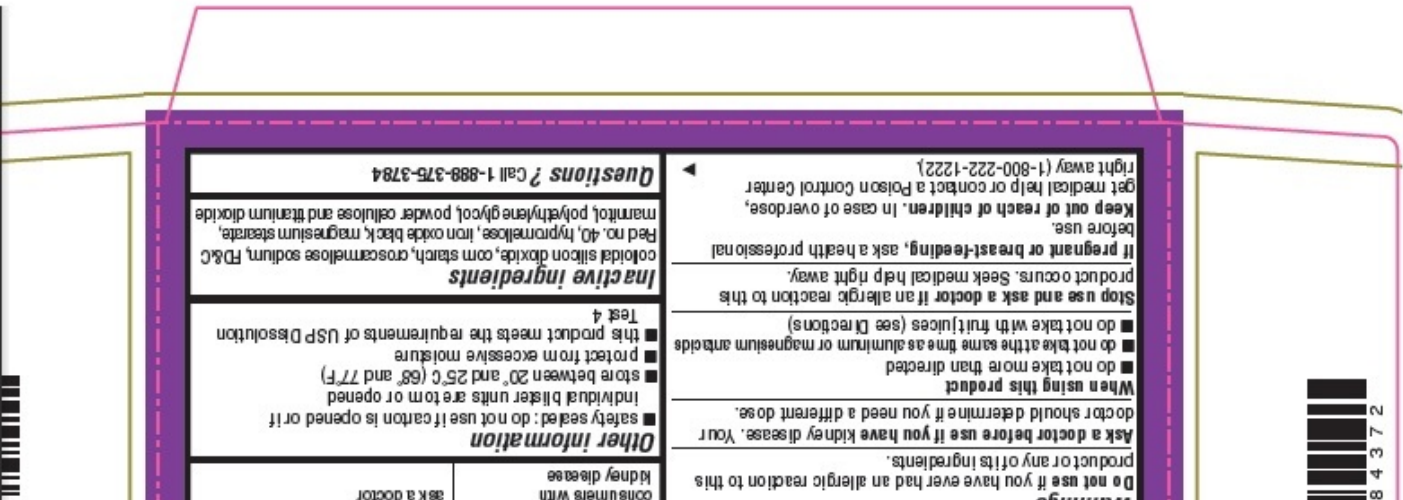
Inactive ingredients

colloidal silicon dioxide, corn starch, croscarmellose sodium, FD&C Red no. 40, hypromellose, iron oxide black, magnesium stearate, mannitol, polyethylene glycol, powder cellulose and titanium dioxide

Questions?

Call 1-888-375-3784

PACKAGE LABEL PRINCIPAL DISPLAY PANEL SECTION



Drug Facts
Active ingredient (in each tablet)
Fexofenadine HCl USP, 60 mg
Purpose
Antihistamine

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

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children under 12 years of age
do not use
ask a doctor
adults 65 years of age and older

TAMPER EVIDENT: INDIVIDUAL BUSTER UNIT SEALED FOR YOUR PROTECTION.
DO NOT USE IF INDIVIDUAL BUSTER UNIT IS OPEN OR TORN.
IMPORTANT: Read the directions and warnings before use.
Keep the carton. It contains important information.



Non-Drowsy
Allergy Relief
Fexofenadine Hydrochloride Tablets, USP
60 mg
Antihistamine



NDC 63868-819-12

Compare to the active ingredient in Allegra® Allergy 12 Hour Tablets*

Non-Drowsy
Allergy Relief
Fexofenadine Hydrochloride Tablets, USP
60 mg
Antihistamine

INDOOR/OUTDOOR ALLERGY RELIEF

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

12 HR



12 Tablets

*This product is not manufactured or distributed by Chatham, Inc. (part of the SanofiGroup), distributor of Allegra® Allergy 12 Hour Tablets. Allegra® is a registered trademark of Aventis sub LLC.

100% SATISFACTION GUARANTEED
Distributed by: C.D.M.A., Inc.®
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www.qualitychoice.com
Questions: 800-935-2362



LOT EXP

ALLERGY RELIEF

fexofenadine hydrochloride 60 mg tablet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63868-819(NDC:55111-783)
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
magnesium stearate (UNII: 70097M6I30)	
mannitol (UNII: 3OWL53L36A)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	
FERROSOFERRIC OXIDE (UNII: XM0M87F357)	
polyethylene glycol 400 (UNII: B697894SGQ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	PINK	Score	no score
Shape	OVAL	Size	5mm
Flavor		Imprint Code	193;R
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63868-819-12	2 in 1 CARTON	05/17/2021	
1		6 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	ANDA076502	05/17/2021	

