ALLEGRA ALLERGY- fexofenadine hydrochloride tablet, film coated
Lil' Drug Store Products, Inc

Allegra ® Allergy

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.

Directions

adults and	

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

- safety sealed: do not use if carton is opened or if individual blister units are torn or opened
- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide blends, magnesium stearate, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch, titanium dioxide

Questions or comments?

call toll-free **1-800-633-1610** or www.allegra.com

Product repackaged and distributed by:
Lil' Drug Store Products, Inc.
9300 Earhart Lane SW
Cedar Rapids, IA 52404

PRINCIPAL DISPLAY PANEL - 5 Tablet Blister Pack Carton

NDC 66715-9707-8
NON-DROWSY

Allegra ®
ALLERGY
fexofenadine HCl tablet
180 mg/antihistamine
24 HR

INDOOR / OUTDOOR ALLERGY RELIEF

✓ Sneezing
✓ Runny Nose

✓ Itchy, Watery Eyes
✓ Itchy Nose or Throat

Actual Size

5
Tablets

Lil'
DrugStore ®

NON-DROWSY

Allegra®
ALLERGY
fexofenadine HCl tablet
180 mg/antihistamine 24HR

Allegra®
ALLERGY 24HR

The makers of Allegra® do not make store brand products. The trade dress of this Allegra® package is subject to trademark protection.

Product manufactured for:
Chattem, Inc., a Sanofi Company
Chattanooga, TN 37409-0219 ©2017
Origin Germany

Product repackaged and distributed by:
Lil' Drug Store Products, Inc.
9300 Earhart Lane SW
Cedar Rapids, IA 52404

NDC 66715-9707-8

NON-DROWSY

Allegra®
ALLERGY
fexofenadine HCl tablet
180 mg/antihistamine 24HR

INDOOR / OUTDOOR ALLERGY RELIEF
- ✓ Sneezing
- ✓ Runny Nose
- ✓ Itchy, Watery Eyes
- ✓ Itchy Nose or Throat

Actual Size

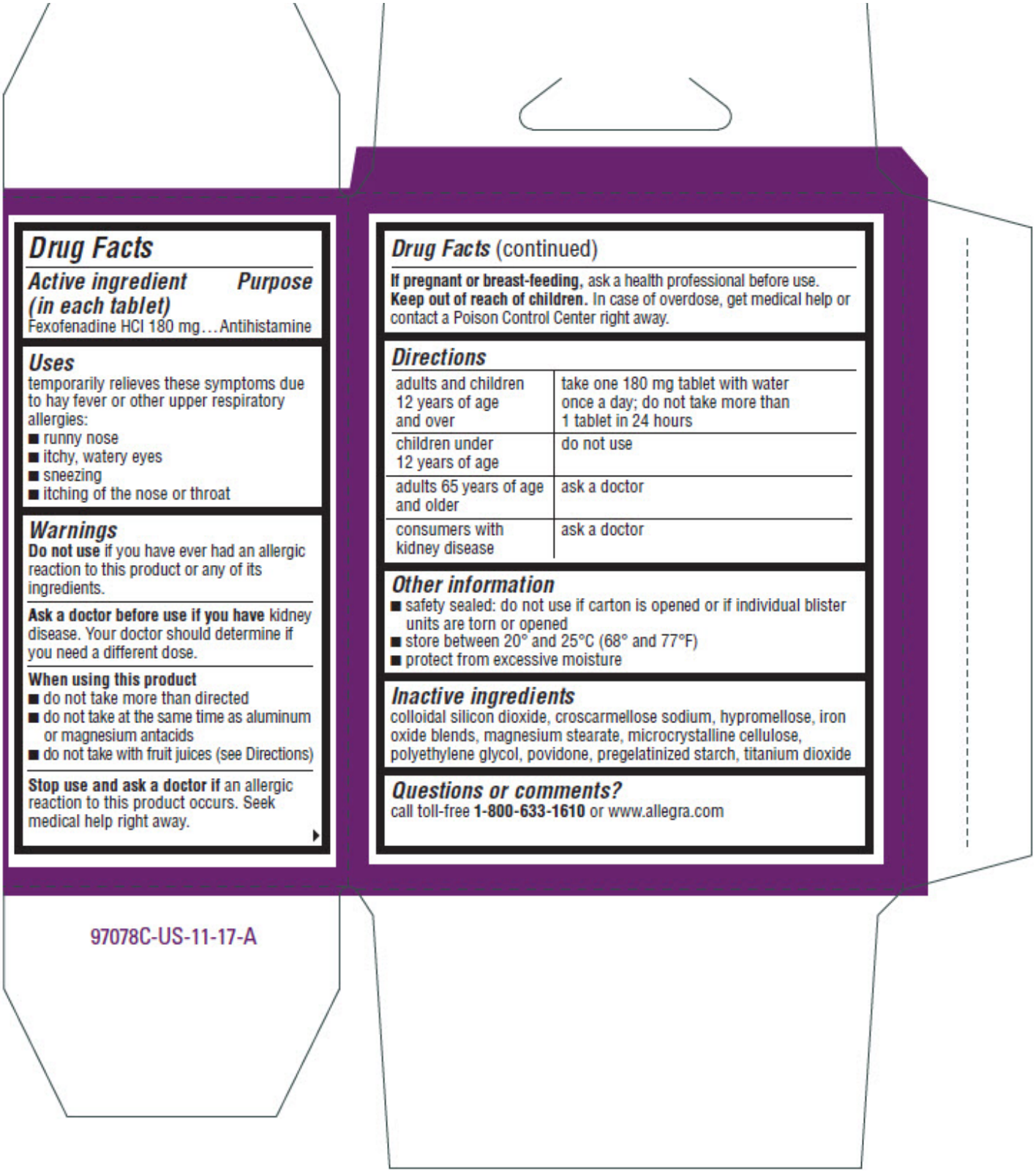
5
Tablets

Lil' DrugStore®

3 66715 97078 7

LOT

EXP



Drug Facts

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Fexofenadine HCl 180 mg… Antihistamine

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Drug Facts (continued)

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97078C-US-11-17-A

ALLEGRA ALLERGY

fexofenadine hydrochloride tablet, film coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66715-9707
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
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POVIDONE, UNSPECIFIED (UNII: FZ989GH94E)	
STARCH, CORN (UNII: O8232NY3SJ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	orange (Peach)	**Score**	no score
Shape	OVAL	**Size**	17mm
Flavor		**Imprint Code**	018;E
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66715-9707-1	1 in 1 CARTON	11/01/2018	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:66715-9707-2	2 in 1 CARTON	11/01/2018	
2		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
3	NDC:66715-9707-8	5 in 1 CARTON	11/01/2018	
3		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
NDA	NDA020872	09/01/2014	

Labeler - Lil' Drug Store Products, Inc (093103646)

Revised: 12/2022

Lil' Drug Store Products, Inc