ALLEGRA ALLERGY- fexofenadine hydrochloride tablet, coated
Chattem, Inc.

Allegra Allergy

Allegra Allergy® Gelcaps – 24 HOUR

Drug Facts

Active ingredient

(in each tablet)

Fexofenadine HCI 180 mg

Purpose

Antihistamine

Uses

temporarily relieves these symptoms due to hay fever or other upper respiratory allergies:

- runny nose
- sneezing
- itchy, water 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.

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 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

croscarmellose sodium, D&C red 28, D&C red 33, FD&C blue 1, gelatin, hydroxypropylcellulose, hydroxypropyl methylcellulose, magnesium stearate, microcrystalline cellulose, PEG-135, pharmaceutical ink, pregelatinized starch, titanium dioxide

Questions or comments?

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

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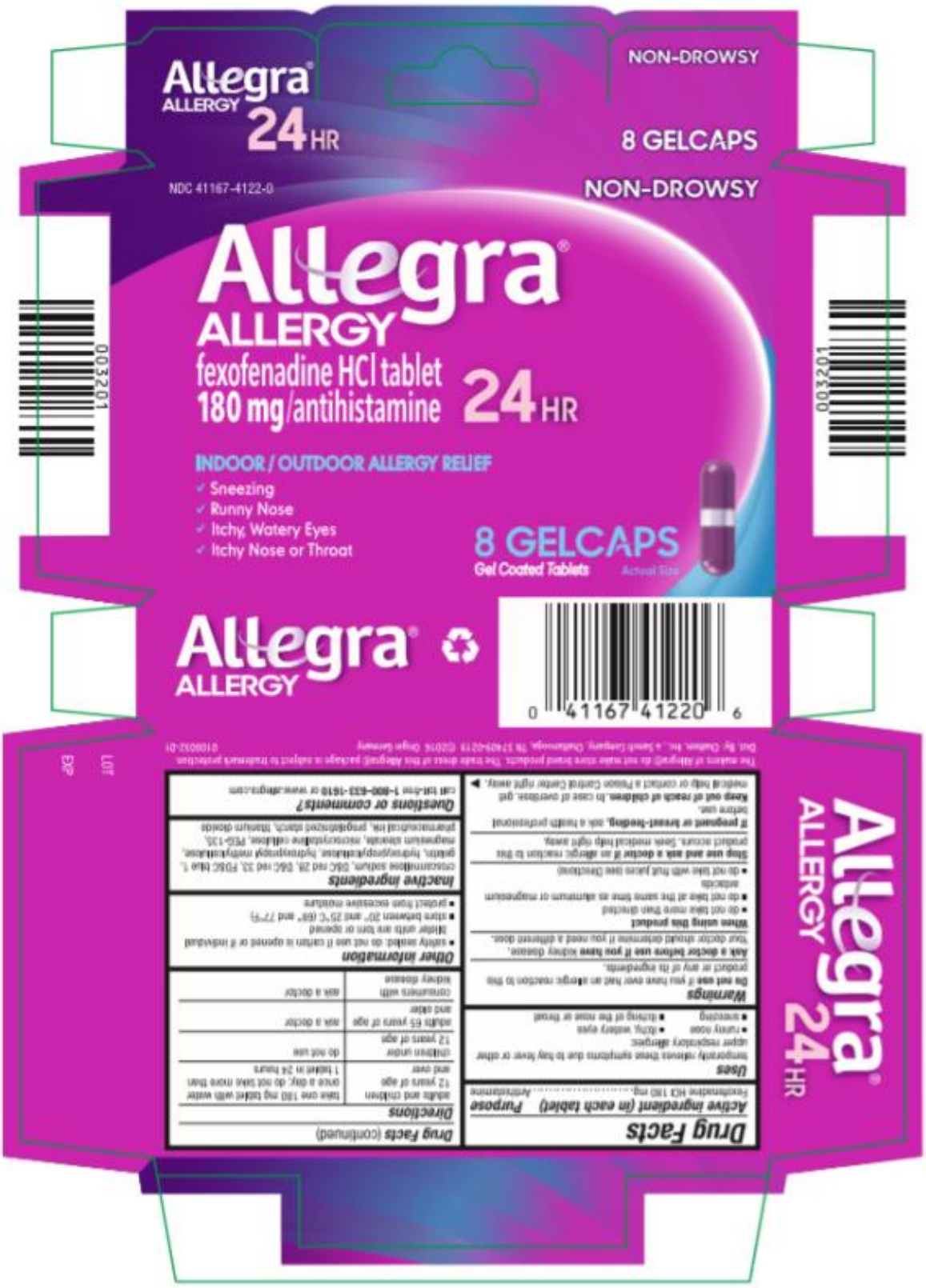
PRINCIPAL DISPLAY PANEL

NDC 41167-4122-0
Allegra® Allergy
Fexofenadine HCl tablets
180 mg/antihistamine
24 HR
8 Gelcaps



PRINCIPAL DISPLAY PANEL

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Fexofenadine HCl tablets
180 mg/antihistamine
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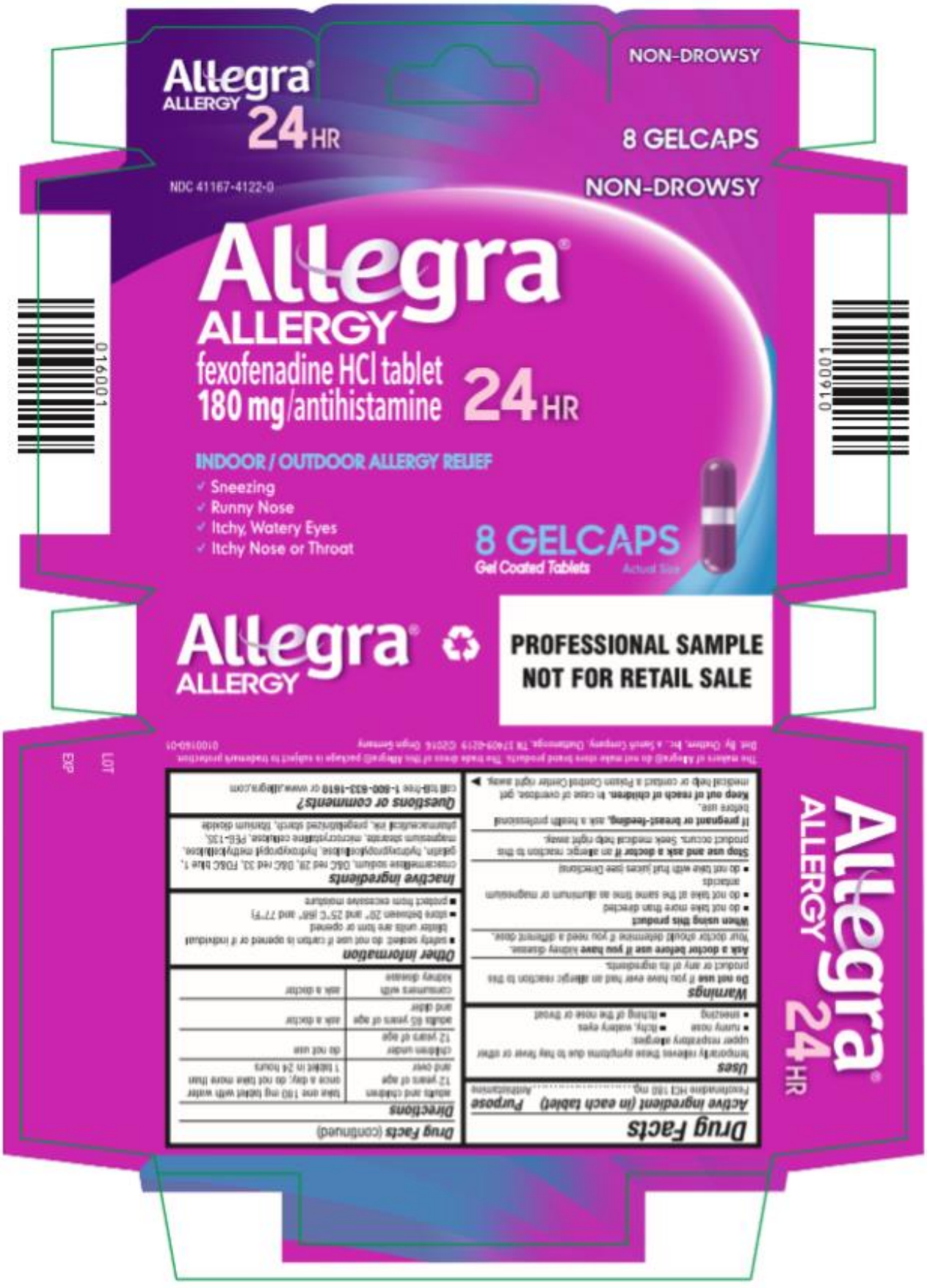
PROFESSIONAL SAMPLE
NOT FOR RETAIL SALE



ALLEGRA ALLERGY

fexofenadine hydrochloride tablet, coated

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:41167-4122
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**
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
D&C RED NO. 28 (UNII: 767IP0Y5NH)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GELATIN (UNII: 2G86QN327L)	
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL 6000 (UNII: 30IQX730WE)	
STARCH, CORN (UNII: O8232NY3SJ)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics			
Color	PURPLE (White band in the middle)	**Score**	no score
Shape	OVAL (Caplet)	**Size**	20mm
Flavor		**Imprint Code**	AG;AG
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:41167-4122-0	1 in 1 CARTON	12/17/2014	
1		8 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:41167-4122-2	1 in 1 CARTON	12/17/2014	
2		60 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:41167-4122-5	1 in 1 CARTON	12/17/2014	
3		24 in 1 BOTTLE; Type 0: Not a Combination Product		
4	NDC:41167-4122-6	1 in 1 CARTON	12/17/2014	
4		32 in 1 BOTTLE; Type 0: Not a Combination Product		

5	NDC:41167-4122-9	2 in 1 CARTON	12/17/2014	06/23/2019
5	NDC:41167-4122-7	40 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
NDA	NDA020872	12/17/2014	

Labeler - Chattem, Inc. (003336013)

Revised: 9/2019 Chattem, Inc.