ALLERGY RELIEF- fexofenadine hydrochloride tablet, film coated
Publix Super Markets Inc

Public Super Markets, Inc. Allergy Relief 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 (1-800-222-1222).

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

- do not use if printed blister unit is broken or torn
- store between 68° -77°F (20° -25°C)
- protect from excessive moisture
- this product meets the requirements of USP Dissolution Test 2

Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, FD&C blue #2 aluminum lake, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, pregelatinized starch, talc, titanium dioxide

Package/Label Principal Display Panel

P

24-HR

allergy relief

FEXOFENADINE HYDROCHLORIDE TABLETS, 180 mg

ANTIHISTAMINE

Indoor/outdoor allergy relief

• Sneezing

• Runny nose

• Itchy, watery eyes

• Itchy nose or throat

NON-DROWSY

ACTUAL SIZE

15 TABLETS

Compare to Allegra® Allergy active ingredient

24-HR
allergyrelief
FEXOFENADINE HYDROCHLORIDE TABLETS, 180 mg
ANTIHISTAMINE

NDC 56062-847-22

24-HR
allergyrelief
FEXOFENADINE HYDROCHLORIDE TABLETS, 180 mg
ANTIHISTAMINE

Indoor/outdoor allergy relief
- Sneezing
- Runny nose
- Itchy, watery eyes
- Itchy nose or throat

NON-DROWSY

ACTUAL SIZE

15 TABLETS

Compare to Allegra® Allergy active ingredient*

84722 63 C1

OPEN OTHER END

CONVENIENT RECLOSING TAB

Drug Facts

Active ingredient (in each tablet) **Purpose**
Fexofenadine HCl 180 mg ..... 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 (1-800-222-1222).

Drug Facts (continued)

Directions

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Inactive ingredients
colloidal silicon dioxide, croscarmellose sodium, FD&C blue #2 aluminum lake, FD&C red #40 aluminum lake, FD&C yellow #6 aluminum lake, lactose monohydrate, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polyvinyl alcohol, pregelatinized starch, talc, titanium dioxide

*This product is not manufactured or distributed by the owner of the registered trademark Allegra®.

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ALLERGY RELIEF

fexofenadine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:56062-847
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)	
FD&C BLUE NO. 2 (UNII: L06K8R7DQK)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	PINK	**Score**	no score
Shape	OVAL	**Size**	18mm
Flavor		**Imprint Code**	L847
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56062-847-22	15 in 1 CARTON	01/14/2022	
1		1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
2	NDC:56062-847-39	1 in 1 CARTON	01/21/2022	

2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:56062-847-95	1 in 1 CARTON	01/21/2022	
3		45 in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA212971	01/14/2022	

Labeler - Publix Super Markets Inc (006922009)

Revised: 1/2022 Publix Super Markets Inc