

**ALLERGY RELIEF- fexofenadine hydrochloride tablet, film coated  
Meijer, Inc.**

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**Meijer Distribution, 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 20° -25°C (68° -77°F)
- protect from excessive moisture
- this product meets the requirements of USP *DissolutionTest 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

## Questions or comments?

**1-800-719-9260**

## Package/Label Principal Display Panel

Compare to Allegra® Allergy 24 HR active ingredient

NON-DROWSY

allergy relief

Fexofenadine Hydrochloride Tablets, 180 mg

Antihistamine

24 HR

INDOOR/OUTDOOR ALLERGY RELIEF

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

15 Tablets

actual size





DST. BY MEIJER  
GRAND RAPIDS, MI  
49544  
www.meijer.com



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

PID 2884376

**Drug Facts**

**Active ingredient (in each tablet)** Fexofenadine HCl 180 mg, Arhistamine

**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, you're on a drug that affects the same time as aluminum or magnesium antacids, or you're taking more than directed.  
 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 breastfeeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact Poison Control. Center right away. (1-800-222-1222)

**Other information**  
 ■ do not use if printed blister unit is broken or torn  
 ■ store between 20°-25° C (68°-77° F)  
 ■ 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, indigo carmine, FD&C yellow #6, polyethylene glycol, polyvinyl alcohol, povidone, starch, talc, titanium dioxide

**Questions or comments?** 1-800-719-9260

**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

**Drug Facts (continued)**

84722 6E C1

**NON-DROWSY**  
**allergy relief**  
 Fexofenadine Hydrochloride Tablets, **180 mg**  
 Antihistamine

**24 HR**

**meijer**

NDC 79481-0847-0  
 Compare to Allegra® Allergy 24 HR active ingredient\*

**NON-DROWSY**  
**allergy relief**  
 Fexofenadine Hydrochloride Tablets, **180 mg**  
 Antihistamine

**24 HR**  
**INDOOR/OUTDOOR ALLERGY RELIEF**  
 Sneezing; Runny Nose; Itchy, Watery Eyes; Itchy Nose or Throat

**15 Tablets** |  actual size

## ALLERGY RELIEF

fexofenadine hydrochloride tablet, film coated

### Product Information

|                                |                |                           |                |
|--------------------------------|----------------|---------------------------|----------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:79481-0847 |
| <b>Route of Administration</b> | ORAL           |                           |                |

### Active Ingredient/Active Moiety

| Ingredient Name   | Basis of Strength          | Strength |
|---|----------------------------|----------|
| <b>FEXOFENADINE HYDROCHLORIDE</b> (UNII: 2S068B75ZU) (FEXOFENADINE - UNII:E6582LOH6V) | FEXOFENADINE HYDROCHLORIDE | 180 mg   |

### Inactive Ingredients

| Ingredient Name  | Strength |
|--|----------|
| <b>CROSCARMELOSE SODIUM</b> (UNII: M28OL1HH48)             |          |
| <b>FD&amp;C BLUE NO. 2</b> (UNII: L06K8R7DQK)              |          |
| <b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)              |          |
| <b>FD&amp;C YELLOW NO. 6</b> (UNII: H77VEI93A8)            |          |
| <b>LACTOSE MONOHYDRATE</b> (UNII: EWQ57Q8I5X)              |          |
| <b>MAGNESIUM STEARATE</b> (UNII: 70097M6I30)               |          |
| <b>MICROCRYSTALLINE CELLULOSE</b> (UNII: OP1R32D61U)       |          |
| <b>POLYETHYLENE GLYCOL, UNSPECIFIED</b> (UNII: 3WJQ0SDW1A) |          |
| <b>POLYVINYL ALCOHOL, UNSPECIFIED</b> (UNII: 532B59J990)   |          |
| <b>TALC</b> (UNII: 7SEV7J4R1U)                             |          |
| <b>TITANIUM DIOXIDE</b> (UNII: 15FIX9V2JP)                 |          |

### Product Characteristics

|                 |      |                     |          |
|-----------------|------|---------------------|----------|
| <b>Color</b>    | PINK | <b>Score</b>        | no score |
| <b>Shape</b>    | OVAL | <b>Size</b>         | 18mm     |
| <b>Flavor</b>   |      | <b>Imprint Code</b> | L847     |
| <b>Contains</b> |      |                     |          |

### Packaging

| # | Item Code        | Package Description                               | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:79481-0847-1 | 1 in 1 CARTON                                     | 10/06/2021           |                    |
| 1 |                  | 30 in 1 BOTTLE; Type 0: Not a Combination Product |                      |                    |
| 2 | NDC:79481-0847-0 | 15 in 1 CARTON                                    | 10/18/2021           |                    |

|   |                  |  |            |
|---|------------------|--|------------|
| 2 |                  | 1 in 1 BLISTER PACK; Type 0: Not a Combination Product |            |
| 3 | NDC:79481-0847-2 | 1 in 1 CARTON  | 10/18/2021 |
| 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                               | 10/06/2021           |                    |

**Labeler** - Meijer, Inc. (006959555)

Revised: 10/2021

Meijer, Inc.