# **FEXOFENADINE HCL-** fexofenadine hcl tablet Walmart Inc.

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Fexofenadine Hydrochloride Tablets USP, 180 mg

#### ACTIVE INGREDIENT(S), in each tablet

Fexofenadine hydrochloride USP, 180 mg

#### **PURPOSE**

**Antihistamine** 

#### USE(S)

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

- § runny nose
- § sneezing
- § itchy, watery 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**

60 mg

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		

#### 180 mg

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 printed foil inner seal on the bottle is torn or missing.
- ☐ store between 20° and 25°C (68° and 77°F)
- □ protect from excessive moisture

#### **INACTIVE INGREDIENTS**

Colloidal silicon dioxide, hypromellose, light liquid paraffin, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, pregelatinized starch, red iron oxide, sodium starch glycolate, talc, titanium dioxide and yellow iron oxide.

#### **QUESTIONS OR COMMENTS**

call toll-free weekdays 9 AM to 5 PM EST at 1-888-588-1418

### PRINCIPAL DISPLAY PANEL

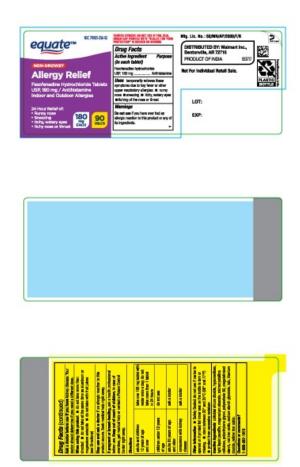
Fexofenadine Hydrochloride Tablets, USP 180 mg 45ct CTN



Fexofenadine Hydrochloride Tablets, USP 180 mg 45ct LBL



Fexofenadine Hydrochloride Tablets, USP 180 mg 90ct LBL



Fexofenadine Hydrochloride Tablets, USP 180 mg 180ct CTN



Fexofenadine Hydrochloride Tablets, USP 180 mg 300ct LBL





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## **FEXOFENADINE HCL**

fexofenadine hcl tablet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79903-358(NDC:69230-300)
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)	
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ8WG20P6)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYSORBATE 80 (UNII: 60ZP39ZG8H)	
STARCH, CORN (UNII: O8232NY3SJ)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	
FERRIC OXIDE YELLOW (UNII: EX43802MRT)	
<b>SODIUM STARCH GLYCOLATE TYPE A POTATO</b> (UNII: 5856J3G2A2)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
POLYETHYLENE GLYCOL 6000 (UNII: 30IQX730WE)	

Product Characteristics						
Color	pink	Score	no score			
Shape	CAPSULE	Size	18mm			
Flavor		Imprint Code	J;44			
Contains						

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-358- 45	1 in 1 CARTON	05/19/2025	
1		45 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:79903-358- 90	2 in 1 CARTON	05/19/2025	
2		90 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:79903-358- 30	300 in 1 BOTTLE; Type 0: Not a Combination Product	05/19/2025	

Marketing Information							
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
ANDA	ANDA204097	05/19/2025					

## Labeler - Walmart Inc. (051957769)

Establishment			
Name	Address	ID/FEI	<b>Business Operations</b>
Legacy Pharmaceutical Packaging, LLC		143213275	relabel(79903-358), repack(79903-358)

Revised: 5/2025 Walmart Inc.