## ALLERGY RELIEF- fexofenadine hcl tablet, film coated Bi-Mart

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## 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
- 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 (1-800-222-1222) right away.

#### **Directions**

adults and children 12 years	take one 180mg tablet with water once a day; do not
of age and over	take more than 1 tablet in 24 hours
children under 12 years of age	do not use
adults 65 years of age and	ask a doctor
older	
consumers with kidney	ask a doctor
disease	

#### Other information

- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture
- each tablet contains: sodium 8 mg
- this product meets the requirements of USP Dissolution Test 2

### Inactive ingredients

anhydrous lactose, colloidal silicon dioxide, corn starch, croscarmellose sodium, hypromellose, lactose monohydrate, polyethylene glycol, pregelatinized starch(maize), red iron oxide, stearic acid, titanium dioxide, yellow iron oxide.

## **Questions?**

Call toll-free **1-844-912-4012** 

NDC 37835-601-09

# COMPARE TO THE ACTIVE INGREDIENT IN ALLEGRA ®ALLERGY 24 HOUR TABLETS\*

**NON-DROWSY** 

#### **ALLERGY RELIEF**

FEXOFENADINE HYDROCHLORIDE 180 mg

24 HOUR ALLERGY

ORIGINAL PRESCRIPTION STRENGTH

**RELIEF OF:** 

**SNEEZING** 

**RUNNY NOSE** 

ITCHY, WATER EYES

ITCHY NOSE OR THROAT

**ANTIHISTAMINE** 

#### 90 Tablets

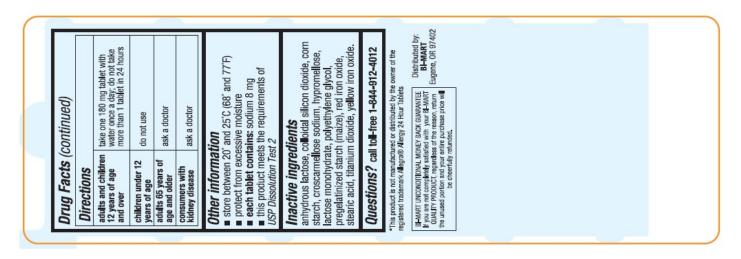
\*This product is not manufactured or distributed by the owner of the registered trademark Allegra <sup>®</sup> Allergy 24 Hour Tablets.

Distributed by:

#### **BI-MART**

Eugene, OR 97402





#### NDC 37835-601-45

COMPARE TO THE ACTIVE INGREDIENT IN ALLEGRA ®ALLERGY 24 HOUR TABLETS\*

**NON-DROWSY** 

**ALLERGY RELIEF** 

FEXOFENADINE HYDROCHLORIDE 180 mg

24 HOUR ALLERGY

ORIGINAL PRESCRIPTION STRENGTH

**RELIEF OF:** 

**SNEEZING** 

RUNNY NOSE

ITCHY, WATER EYES

ITCHY NOSE OR THROAT

ANTIHISTAMINE

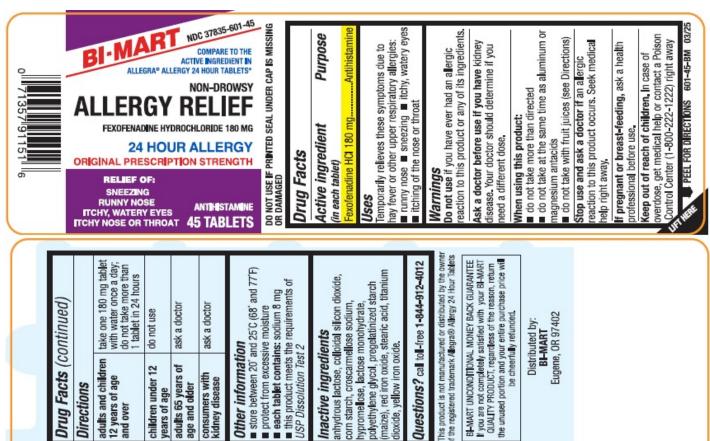
#### 45 Tablets

\*This product is not manufactured or distributed by the owner of the registered trademark Allegra <sup>®</sup> Allergy 24 Hour Tablets.

Distributed by:

#### **BI-MART**

Eugene, OR 97402



#### **ALLERGY RELIEF**

fexofenadine hcl tablet, film coated

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37835-601	
Route of Administration	ORAL			

## **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
	FEXOFENADINE HYDROCHLORIDE	180 mg

Inactive Ingredients			
Ingredient Name	Strength		
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
STARCH, CORN (UNII: O8232NY3SJ)			
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)			
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
STEARIC ACID (UNII: 4ELV7Z65AP)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			
FERRIC OXIDE RED (UNII: 1K09F3G675)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
FERRIC OXIDE YELLOW (UNII: EX43802MRT)			

Product Characteristics				
Color pink Score no score				
Shape	OVAL	Size	18mm	
Flavor		Imprint Code	SG202	
Contains				

F	Packaging					
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:37835-601- 45	45 in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2025			
2	NDC:37835-601- 09	90 in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2025			

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
ANDA	ANDA204507	09/16/2015		

## **Labeler -** Bi-Mart (027630078)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
ScieGen Pharmaceuticals, INC.		079391286	manufacture(37835-601)	

Revised: 4/2025 Bi-Mart