

ALLERGY RELIEF- fexofenadine hydrochloride tablet
YYBA CORP

YYBA (as PLD) - WELMATE ANTIHISTAMINE ALLERGY RELIEF (73581-011)

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 of age and over	take one 180 mg tablet with water once a day; do not take more than 1 tablets in 24 hours
children under 12 years of age	

children under 12 years of age	do not use
adults 65 years and older	ask a doctor
consumers with kidney disease	ask a doctor

Other information

- store between 20° and 25°C (68° and 77°F)
- protect from excessive moisture
- **each tablet contains:**sodium 8.2 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 1-800-935-6737



DO NOT USE IF PRINTED SEAL UNDER CAP IS MISSING OR DAMAGED

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LIFT HERE

PEEL FOR DIRECTIONS G7022-100-103-0

Drug Facts (continued)

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Questions? call toll-free 1-866-933-6337

*This product is not manufactured or distributed by the owner of the registered trademark Allegra® Allergy 24 Hour Tablets



Distributed by:
WellSpring
Albion, NY 10952, U.S.A.



Why pay more?

wellspringmeds.com



3 73581 00062 3

Made in the USA





Compare to the active ingredient in Allegra® Allergy 24HR Tablets*

ANTIHISTAMINE

Allergy Relief

Fexofenadine Hydrochloride **180 MG**

Non-drowsy 24 hour allergy relief of:

Sneezing
Runny nose
Itchy watery eyes
Itchy nose or throat

Gluten Free
ORIGINAL PRESCRIPTION STRENGTH

40 TABLETS

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PEEL FOR DIRECTIONS

LIFT HERE

ANTIHISTAMINE

Allergy Relief

Made in the USA



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Wellspring
Airmont, NY 10952, U.S.A.

Why pay more?
wellspringmeds.com



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ALLERGY RELIEF			
fexofenadine hydrochloride tablet			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73581-011
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		
ANHYDROUS LACTOSE (UNII: 3SY5LH9PMK)			
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)			
STARCH, CORN (UNII: O8232NY3SJ)			
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)			
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)			
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)			
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)			

CORN (UNII: 0N8672707O)

FERRIC OXIDE RED (UNII: 1K09F3G675)

STEARIC ACID (UNII: 4ELV7Z65AP)

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

FERRIC OXIDE YELLOW (UNII: EX438O2MRT)

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73581-011-80	100 in 1 BOTTLE; Type 0: Not a Combination Product	03/25/2021	
2	NDC:73581-011-40	40 in 1 BOTTLE; Type 0: Not a Combination Product	01/02/2025	
3	NDC:73581-011-10	10 in 1 BOTTLE; Type 0: Not a Combination Product	11/27/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA204507	03/25/2021	

Labeler - YYBA CORP (006339772)

Revised: 11/2025

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