FEXOFENADINE HCL- fexofenadine hcl tablet Meijer, Inc.
Fexofenadine HCl Tablets, USP 180mg
Antihistamine
INDOOR / OUTDOOR ALLERGY RELIEF: Sneezing Runny Nose Itchy, Watery Eyes Itchy Nose or Throat
Active Ingredient (in each tablet)
Fexofenadine HCl USP, 180mg
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.

Directions

disease

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 ask a doctor and older consumers with kidney ask a doctor

Other information

\square safety sealed: do not use if imprinted foil under bottle cap is opened or	torn
☐ store between 20° and 25°C (68° and 77°F)	
□ protect from excessive moisture	

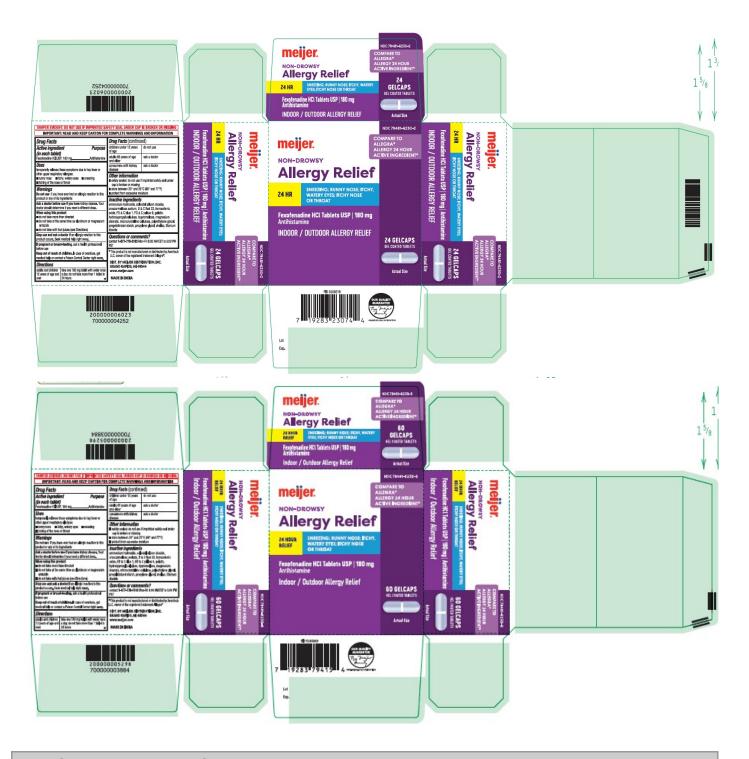
Inactive ingredients

colloidal silicon dioxide, croscarmellose sodium, hypromellose, iron oxide black, iron oxide red, iron oxide yellow, lactose monohydrate, macrogol, magnesium stearate, microcrystalline cellulose, pregelatinized starch, titanium dioxide

Questions or comments?

contact 1-877-770-3183 Mon-Fri 8:00 AM EST to 5:00 PM PST

Principal Display Panel



FEXOFENADINE HCL

fexofenadine hcl tablet

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (So	ource)	NDC:79481	-6230
Route of Administration	ORAL				
A ation to our disast/A ation	Na :- 4				
Active Ingredient/Active Moiety					
Ingredient Name Basis of Strength Streng				Strength	
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - FEXOFENADINE				190 ma	

UNII:E6582LOH6V) HYDROCHLORIDE

Inactive Ingredients		
Ingredient Name	Strength	
D&C RED NO. 28 (UNII: 767IP0Y5NH)		
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)		
GELATIN (UNII: 2G86QN327L)		
HYPROMELLOSES (UNII: 3NXW29V3WO)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
D&C RED NO. 33 (UNII: 9DBA0SBB0L)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		
CROSCARMELLOSE SODIUM (UNII: M280L1HH48)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
STARCH, CORN (UNII: O8232NY3SJ)		
POLYETHYLENE GLYCOL 6000 (UNII: 30IQX730WE)		
HYDROXYPROPYL CELLULOSE (1600000 WAMW) (UNII: RFW2ET671P)		

Product Characteristics			
Color	gray (White band in the middle)	Score	no score
Shape	OVAL (capsule shaped tablet)	Size	20mm
Flavor		Imprint Code	G18
Contains			

F	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:79481- 6230-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2024		
2	NDC:79481- 6230-2	24 in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2025		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA211075	07/01/2024	

Labeler - Meijer, Inc. (006959555)

Revised: 12/2025 Meijer, Inc.