EQUATE ALLERGY RELIEF- fexofenadine hydrochloride tablet Walmart Inc.

Fexofenadine Hydrochloride Tablets USP, 180 mg

Active ingredient(s)

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

Ask a doctor or pharmacist 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

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

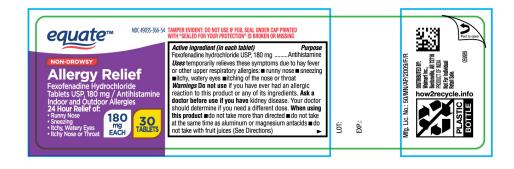
Colloidal silicon dioxide, hypromellose, light liquid paraffin, magnesium stearate, microcrystalline cellulose, polyethylene glycol, polysorbate 80, pregelatinized starch, red iron oxide, sodium starch glycolate, talc, titanium dioxide and yellow iron oxide.

Questions

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

Principal Display Panel

Fexofenadine Hydrochloride Tablets, USP 180 mg 30 ct Container Label





Drug Facts (Continued) Stop use and ask a doctor if an allergic reaction to professional before use. Keep out of reach of ch Directions	o this product occurs. Seek medical help right away. If pregnant or breast-feeding, ask a health Ildren. In case of overdose, get medical help or contact a Poison Control Center right away.
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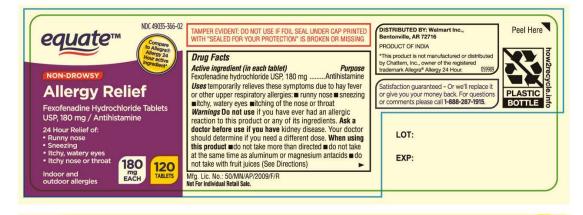
Fexofenadine Hydrochloride Tablets, USP 180 mg 30ct Carton Label



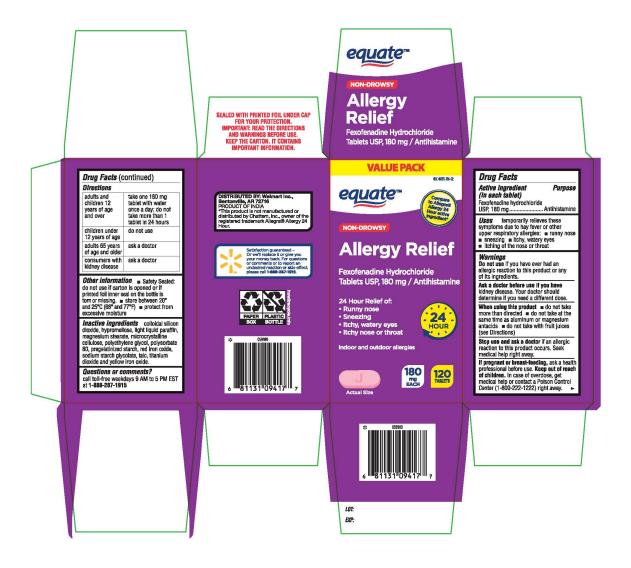
Fexofenadine Hydrochloride Tablets, USP 180 mg 2x30 ct Carton Label



Fexofenadine Hydrochloride Tablets, USP 180 mg 120 ct Carton Label



	ils product occurs. Seek medical help right away. If pregnant or breast-feeding, ask a health profession 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 tablet in 24 hours
children under 12 years of age	do not use
adults 65 years of age and older	ask a doctor
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EQUATE ALLERGY RELIEF

fexofenadine hydrochloride tablet

Product Information

Product Type

Ingredient Name	Basis of Strengt	h Strength
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADIN UNII:E6582LOH6V)	E - FEXOFENADINE HYDROCHLORIDE	180 mg
Inactive Ingredients		
Ingredient Name		Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)		
HYPROMELLOSE 2910 (6 MPA.S) (UNII: 0WZ 8WG20P6)		
LIGHT MINERAL OIL (UNII: N6K5787QVP)		
MAGNESIUM STEARATE (UNII: 70097M6I30)		
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)		
POLYSORBATE 80 (UNII: 60ZP39ZG8H)		
STARCH, CORN (UNII: 08232NY3SJ)		
FERRIC OXIDE RED (UNII: 1K09F3G675)		
FERRIC OXIDE YELLOW (UNII: EX43802MRT)		
SODIUM STARCH GLYCOLATE TYPE A POTATO (UNII: 5856J3G2A2)		
TALC (UNII: 7SEV7J4R1U)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

Product Characteristics

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

Packaging

#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-366- 54	1 in 1 CARTON	04/14/2011	
1		30 in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:49035-366- 59	2 in 1 CARTON	04/14/2011	
2		30 in 1 BOTTLE; Type 0: Not a Combination Product		
3	NDC:49035-366- 02	1 in 1 CARTON	04/14/2011	
3		120 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	ANDA204097	04/14/2011	

Labeler - Walmart Inc. (051957769)

Establishment

Name	Address	ID/FEI	Business Operations
Legacy Pharmaceutical Packaging, LLC		143213275	repack(49035-366) , relabel(49035-366)

Revised: 5/2023

Walmart Inc.