FEXOFENADINE HCL- fexofenadine hcl tablet Preferred Pharmaceuticals, Inc.

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.
- I 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

NDC 68788-7449

Repackaged by Preferred Pharmaceuticals, Inc.

Fexofenadine HCI Tablets 180mg	PREFERRED Pharmaceuticals, Inc.	CAUTION: Federal law PROHIBITS transfer of this drug to any person other thean the patient for whom it was prescribed	Fexofenadine HCl Tablets 180mg Qty: Ins: Lo#: Bat#: Prod# (NDC):	Log
Generic for Allegra Active ingredient (in each tablet): Fexofenadine HCl 180mgAntihistamine Pkg Size: Exp Date: Lot#: Batch#:	=		Fexofenadine HCl Tablets 180mg Qty: Ins: Lot#: Bat#: Prod# (NDC):	Chart
Ins: Ins: Mfg: Camber Consumer Care Prod#: Multimatine Non-Drowy, Store Borward, 20*-25*C (6*-77*), project from year, Store Borward, This product meets the requirements of USP Dissolution Test 7. Do not use if you have sever had an altering reason to before use if you have shaday duesse. When using this product, do bot have another than directed to do not take at the	Directions English tablet(s) hours.	Instrucciones Espanol: tableta(s) horas.	Fexofenadine HCl Tablets 180mg Qty: Insurance NDC: Lot#: Bat#:	Billing
this product or any of its ingredients. Ask if doctor before use of yout have kindipy disease. When using that the same time, as aluminum or magnesium andpacits, do not take with inut juice; soon use and ask a doctor of an each state that the same same same same same high ways. If pregnant dy treast-checking, ask a blashin professional lebore use. Leop out of rack of children -4Hour. Tablet is capsule slapped, punk, imprinted with J 44	Take	Toma cada	Fexofenadine HCl Tablets 180mg Qty: Ins: Lot#: Bat#: Prod# (NDC):	Patient

Fexofenadine Hydrochloride Tablets, USP 180 mg

FEXOFENADINE HCI	_					
fexofenadine hcl tablet	-					
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source	e) NDC:68788-7449(NDC	:69230-300)		
Route of Administration	ORAL					
	NA . 1 . 1					
Active Ingredient/Active	-					
-	edient Name		Basis of Strength	Strength		
FEXOFENADINE HYDROCHLORIDE (UNII: 2S068B75ZU) (FEXOFENADINE - UNII: E6582L0H6V)FEXOFENADINE - HYDROCHLORIDE				180 mg		
Inactive Ingredients						
mactive myreulents	Ingredient N	200		Strength		
SILICON DIOXIDE (UNII: ETJ7Z6X		Strength				
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: 1K09F3	3G675)					
FERRIC OXIDE YELLOW (UNII: EX	(43802MRT)					
SODIUM STARCH GLYCOLATE T	YPE A POTATO (UNI	I: 5856J3G2A2)				
TALC (UNII: 7SEV7J4R1U)						

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)									
PC	DLYETHYLENE G	LYCOL 6	000 (UNII: 30IQX73	OWE)					
-			-						
Product Characteristics									
	olor		PINK		Score			no score	
Sł	nape		CAPSULE		Size			18mm	
Fla	avor				Imprint Cod	е		J;44	
Сс	ontains								
_									
Pa	ackaging								
#	Item Code		Package Description		Marketing Start Date		Marketing End Date		
1	NDC:68788- 7449-3	30 in 1 E Product	BOTTLE; Type 0: Not a Combination		12/27/2019				
2	NDC:68788- 7449-6	60 in 1 E Product	1 BOTTLE; Type 0: Not a Combination ct			12	/27/2019		
3	NDC:68788- 7449-9	90 in 1 E Product	BOTTLE; Type 0: Not a Combination		12/27/2019				
4	NDC:68788- 7449-1	100 in 1 Product	0 in 1 BOTTLE; Type 0: Not a Combination oduct			12/27/2019			
Marketing Information									
	Marketing Category			Marketing Start Date		Marketing End Date			
AN	DA	ANDA204097		08/19/2016					

Labeler - Preferred Pharmaceuticals, Inc. (791119022)

Registrant - Preferred Pharmaceuticals, Inc. (791119022)

Establishment							
Name	Address	ID/FEI	Business Operations				
Preferred Pharmaceuticals, Inc.		791119022	REPACK(68788-7449)				

Revised: 4/2024

Preferred Pharmaceuticals, Inc.