

**FEXOFENADINE HCL- fexofenadine hcl tablet, film coated
NuCare Pharmaceuticals, Inc.**

FEXOFENADINE HYDROCHLORIDE TABLETS USP, 180 mg

Allergy

ALLERGY

Active ingredient (in each film-coated tablet)

Fexofenadine HCl USP, 180 mg

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

adults and children 12 years of age and over	take one 180mg 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 bottle is torn or missing
- 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, lactose monohydrate, pregelatinized starch (maize), stearic acid, opadry pink 03B84893 containing hypromellose, polyethylene glycol, red iron oxide, titanium dioxide and yellow iron oxide.

Questions or comments?

Call 1-888-588-1418

Distributed by:
Camber Consumer Care, Inc.
Piscataway, NJ 08854, USA,

Made in USA

**Camber Consumer Care - Compare to the active ingredient in Allegra®
Allergy 24 Hour Tablets**

Allergy Relief - 24 HOUR

FEXOFENADINE HYDROCHLORIDE TABLETS USP, 180 mg

Antihistamine

Indoor & Outdoor Allergies

NuCare Pharmaceuticals, Inc.

NDC 68071-4568-9
Lot #: 000000 Exp. Date: 00-00

Fexofenadine HCl 180mg
#90 Tablets

Each tablet contains:
Fexofenadine HCl USP, 180mg..... Antihistamine
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.
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.
In case of overdose, get medical help or contact a Poison
Control Center right away (1-800-222-1222).
Oblong Peach Tablet Debossed:
"202" on one side "SG" on the other side

Rev. 01/18

Fexofenadine HCl 180mg
#90 Tablets Exp Date: 00-00
NDC 68071-4568-09 AWP:
Mfg NDC 69230-202-01
Lot #: 000000 Rx # 23140378

Fexofenadine HCl 180mg
#90 Tablets Exp Date: 00-00
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Fexofenadine HCl 180mg
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NDC 68071-4568-09
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Lot #: 000000 Rx # 23140378

Rx # 23140378

Distributed by:
Gambler Consumer Care, Inc. Pleasantway, NJ 08864
P 1349090

Packaged by:
NuCare Pharmaceuticals, Inc.
Orange, CA 92867

Call your doctor for medical advice about side effects.
You may report side effects to FDA at 1-800-FDA-1088.

Patient Instructions:

Take every hours
times a day.

68071456809*90*000000*000000

Product #: P1349090

WARNING: KEEP OUT OF REACH OF CHILDREN. STORE AT CONTROLLED TEMPERATURE 68-77°F.

FEXOFENADINE HCL

fexofenadine hcl tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68071-4568(NDC:69230-202)
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: 3S5Y5LH9PMK)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
STARCH, CORN (UNII: O8232NY3SJ)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
LACTOSE MONOHYDRATE (UNII: EWQ57Q8I5X)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
FERRIC OXIDE RED (UNII: 1K09F3G675)	

TITANIUM DIOXIDE (UNII: 15FIX9V2JP)

FERRIC OXIDE YELLOW (UNII: EX438O2MRT)

Product Characteristics

Color	pink	Score	no score
Shape	CAPSULE	Size	17mm
Flavor		Imprint Code	SG;202
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-4568-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	09/20/2018	
2	NDC:68071-4568-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	09/20/2018	

Marketing Information

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

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals, Inc.		010632300	repack(68071-4568)

Revised: 1/2022

NuCare Pharmaceuticals, Inc.