

IBUPROFEN- ibuprofen tablet, film coated
NuCare Pharmaceuticals, Inc.

IBUPROFEN 400 MG - 600 MG AND 800 MG TABLETS

800 mg (white to off-white, capsule shaped, biconvex, film-coated tablets debossed with 123 on one side and plain on other side)

NDC 68071-5140-3 BOTTLES OF 3

NDC 68071-5140-6 BOTTLES OF 6

NuCare Pharmaceuticals, Inc.

NDC: 68071-5140-3
Ibuprofen 800mg
#3 Tablets
Capsule Shaped White/Off-White Tablet
Debossed: '123' on one side

Each tablet contains:
Ibuprofen, USP 800mg

Warning: Take with food or milk
Product #: P0795003ER
Rx Only

Manufactured by:
Markans Pharma Ltd. Verna, Goa-403 722.
India

Packaged By:
NuCare Pharmaceuticals, Inc. Orange, CA 92667
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.

GTIN 00368071614032
Serial# 000000000002
Exp. Date 00-00
LOT#: 000000

Rev 01/01/19

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

3 6807151403 2
Ibuprofen 800mg
#3 Tablets Serial# 00000000002
Lot: 000000 NDC: 68071-5140-03
Exp.: 00-00 MFR NDC: 49483-804-60

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:68071-5140(NDC:49483-604)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	800 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL (UNII: 532B59J990)	
STARCH, PREGELATINIZED CORN (UNII: O8232NY3SJ)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	Score	no score
Shape	CAPSULE	Size	19mm
Flavor		Imprint Code	123
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-5140-3	3 in 1 BOTTLE; Type 0: Not a Combination Product	12/31/2019	
2	NDC:68071-5140-6	6 in 1 BOTTLE; Type 0: Not a Combination Product	01/06/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	12/30/2015	

Labeler - NuCare Pharmaceuticals, Inc. (010632300)

Establishment

Name	Address	ID/FEI	Business Operations
NuCare Pharmaceuticals, Inc.		010632300	repack(68071-5140)

Revised: 1/2025

NuCare Pharmaceuticals, Inc.