

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

IBUPROFEN 400 MG - 600 MG AND 800 MG TABLETS

ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

HOW SUPPLIED

400mg (white to of white, round, biconvex, film coated tablets debossed with 121 on one side and plain on the other side)

NDC 68071-5154-1 BOTTLES OF 100

400mg Ibuprofen 100 count label

NDC: 68071-5154-1

Ibuprofen 400mg
#100 Tablets

See manufacturer's label
for full list of ingredients.

Product #: R0775100
Rx Only

Ibuprofen 400mg
Lot: 000000 NDC: 68071-5154-01
MFR NDC: 49483-602-01 Exp.: 00-00
Serial# 00000000002

Ibuprofen 400mg
Lot: 000000 NDC: 68071-5154-01
MFR NDC: 49483-602-01 Exp.: 00-00
Serial# 00000000002



GTIN 00368071515411
Serial# 00000000002
Exp. Date 00-00
LOT#: 000000

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

Manufactured by: 3 68071 51541
Marksans Pharma Ltd. Verna,
Goa-403 722, India
Packed By:
NuCare Pharmaceuticals, Inc.
Orange, CA 92867

Take _____ every _____ hours
_____ times a day.

WARNING: KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED TEMPERATURE 68-77°F.

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:68071-5154(NDC:49483-602)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XY110QM) (IBUPROFEN - UNII:WK2XY110QM)	IBUPROFEN	400 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	ROUND	Size	13mm
Flavor		Imprint Code	121
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-5154-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/17/2020	

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	relabel(68071-5154)

Revised: 2/2021

NuCare Pharmaceuticals,Inc.