

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

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**IBUPROFEN 400 MG - 600 MG AND 800 MG TABLETS**

**ibuprofen tablets 400 mg - 600 mg- 800 mg medguide**

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

**NuCare Pharmaceuticals, Inc.**

**Ibuprofen 800mg #100 Tablets**

NDC: 68071-5135-0

Lot: 000000 NDC: 68071-5135-00  
MFR NDC: 49483-604-01 Exp.: 00-00  
Serial# 00000000002

**Ibuprofen 800mg**  
Lot: 000000 NDC: 68071-5135-00  
MFR NDC: 49483-604-01 Exp.: 00-00  
Serial# 00000000002

GTIN 00368071513509  
Serial# 000000000002  
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.

See manufacturer's label for full list of ingredients.

Product #: R0285100  
**Rx Only**

Manufactured by: 3 68071 51350 9  
Marksans Pharma Ltd. Verna,  
Goa-403 722, India

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

**Patient Instructions:**  
Take \_\_\_\_\_ every \_\_\_\_\_ hours  
\_\_\_\_\_ times a day.

Rev 01/01/19

**WARNING: KEEP OUT OF REACH OF CHILDREN**

**STORE AT CONTROLLED TEMPERATURE 68-77°F.**



## IBUPROFEN

ibuprofen tablet, film coated

**Product Information**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:68071-5135(NDC:49483-604)
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	800 mg

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
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**

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	123
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-5135-0	100 in 1 BOTTLE; Type 0: Not a Combination Product	12/16/2019	

**Marketing Information**

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

**Labeler** - NuCare Pharmaceuticals, Inc. (010632300)**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
NuCare Pharmaceuticals, Inc.		010632300	relabel(68071-5135)

