

**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**

**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 68071-5134-6

 NuCare Pharmaceuticals, Inc.

NDC: 68071-5134-6

**Ibuprofen 400mg**

**#6 Tablets**

Round White/Off-White Tablet Debossed:  
'121' on one side

Each tablet contains:  
Ibuprofen, USP 400mg

Warning: Take with food or milk.

Product #: P0793006ER  
Rx Only



3 6807151346 2

Ibuprofen 400mg  
#6 Tablets Serial# 00000000002  
Lot: 000000 NDC: 68071-5134-06  
Exp.: 00-00 MFR NDC: 49483-602-01



OTIN 00388071613482  
Serial# 00000000002  
Exp. Date 00-00  
LOT#: 000000

Rev. 01/01/19

Manufactured by:  
Markasans Pharma Ltd, Verma, 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.

**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:68071-5134(NDC:49483-602)
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>IBUPROFEN</b> (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	400 mg

**Inactive Ingredients**

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

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	no score
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<b>Shape</b>	ROUND	<b>Size</b>	13mm
<b>Flavor</b>		<b>Imprint Code</b>	121
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-5134-6	6 in 1 BOTTLE; Type 0: Not a Combination Product	12/13/2019	

### 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(68071-5134)

Revised: 6/2024

NuCare Pharmaceuticals, Inc.