

**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 68071-5126-3 BOTTLES OF 30

NDC 68071-5126-9 BOTTLES OF 90

NDC: 68071-5126-9  
**Ibuprofen 400mg**  
**#90 Tablets**

**Ibuprofen 400mg**  
 Lot: 000000 NDC: 68071-5126-09  
 MFR NDC: 49483-602-50 Exp.: 00-00  
 Serial# 00000000002

**Ibuprofen 400mg**  
 Lot: 000000 NDC: 68071-5126-09  
 MFR NDC: 49483-602-50 Exp.: 00-00  
 Serial# 00000000002



GTIN 00368071512694  
 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 51269  
 Marksans Pharma Ltd, Verna,  
 Goa-403 722, India  
 Packaged By:  
 NuCare Pharmaceuticals, Inc.  
 Orange, CA 92867  
 Patient Instructions  
 Take \_\_\_\_\_ every \_\_\_\_\_ hours  
 \_\_\_\_\_ times a day.  
 8807151269\*90-000000-000000  
 Rev 01/01/19

Each tablet contains:  
 Ibuprofen, USP 400mg

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

Product #: P0116090  
**Rx Only**

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:68071-5126(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>CROSCARMELLOSE 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 |
| <b>Shape</b> | ROUND | <b>Size</b>  | 13mm     |

| <b>Flavor</b>                |  | <b>Imprint Code</b>                               | 121                  |                    |
|------------------------------|--|---|----------------------|--------------------|
| <b>Contains</b>              |  |   |                      |                    |
| <b>Packaging</b>             |  |   |                      |                    |
| #                            | Item Code                                | Package Description                               | Marketing Start Date | Marketing End Date |
| 1                            | NDC:68071-5126-9                         | 90 in 1 BOTTLE; Type 0: Not a Combination Product | 12/11/2019           |                    |
| 2                            | NDC:68071-5126-3                         | 30 in 1 BOTTLE; Type 0: Not a Combination Product | 12/11/2019           |                    |
| <b>Marketing Information</b> |  |   |                      |                    |
| Marketing Category           | Application Number or Monograph Citation | Marketing Start Date                              | Marketing End Date   |                    |
| ANDA                         | ANDA090796                               | 12/30/2015  |                      |                    |

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

| <b>Establishment</b>         |         |           |                     |
|------------------------------|---------|-----------|---------------------|
| Name                         | Address | ID/FEI    | Business Operations |
| NuCare Pharmaceuticals, Inc. |         | 010632300 | repack(68071-5126)  |

Revised: 6/2024

NuCare Pharmaceuticals, Inc.