

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

**HOW SUPPLIED**

- NDC 68071-4009-7 Bottles of 15
- NDC 68071-4009-2 Botles of 20
- NDC 68071-4009-1 Bottles of 21
- NDC 68071-4009-3 Bottles of 30
- NDC 68071-4009-4 Bottles of 40
- NDC 68071-4009-5 Bottles of 50
- NDC 68071-4009-6 Bottles of 60
- NDC 68071-4009-9 Bottles of 90

**600mg Ibuprofen Package Label**

**NuCare Pharmaceuticals, Inc.**

NDC: 68071-4009-3  
**Ibuprofen 600mg**  
**#30 Tablets**

Each tablet contains  
Ibuprofen, USP 600mg

Capsule Shaped White/Off-White Tablet  
Debossed: '122' on one side

Product #: P0117030  
**Rx Only**

**Ibuprofen 600mg**  
Lot: 000000 NDC: 68071-4009-03  
MFR NDC: 49483-603-50 Exp.: 00-00  
Serial# 00000000002

**Ibuprofen 600mg**  
Lot: 000000 NDC: 68071-4009-03  
MFR NDC: 49483-603-50 Exp.: 00-00  
Serial# 00000000002

GTIN 00368071400939  
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.

Take \_\_\_\_\_ times a day, \_\_\_\_\_ every \_\_\_\_\_ hours

Manufactured by: Marsans Pharma Ltd. Verna, Goa-403 722, India  
Packaged By: NuCare Pharmaceuticals, Inc. Orange, CA 92867

Rev 01/01/19  
**WARNING: KEEP OUT OF REACH OF CHILDREN** **STORE AT CONTROLLED TEMPERATURE 68-77°F.**

<b>IBUPROFEN</b>			
ibuprofen tablet, film coated			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:68071-4009(NDC:49483-603)
<b>Route of Administration</b>	ORAL		
<b>Active Ingredient/Active Moiety</b>			
	<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
	IBUPROFEN (UNII: WK2XYI10 QM) (IBUPROFEN - UNII:WK2XYI10 QM)	IBUPROFEN	600 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	18mm
Flavor		Imprint Code	122
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-4009-7	15 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
2	NDC:68071-4009-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
3	NDC:68071-4009-1	21 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
4	NDC:68071-4009-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
5	NDC:68071-4009-4	40 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
6	NDC:68071-4009-5	50 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
7	NDC:68071-4009-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	
8	NDC:68071-4009-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	07/19/2017	

## 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(68071-4009)