

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 _____ every _____ hours _____ times a day.

Manufactured by: Mansans 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.

IBUPROFEN			
ibuprofen tablet, film coated			
Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:68071-4009(NDC:49483-603)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	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)

Revised: 7/2024

NuCare Pharmaceuticals, Inc.