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

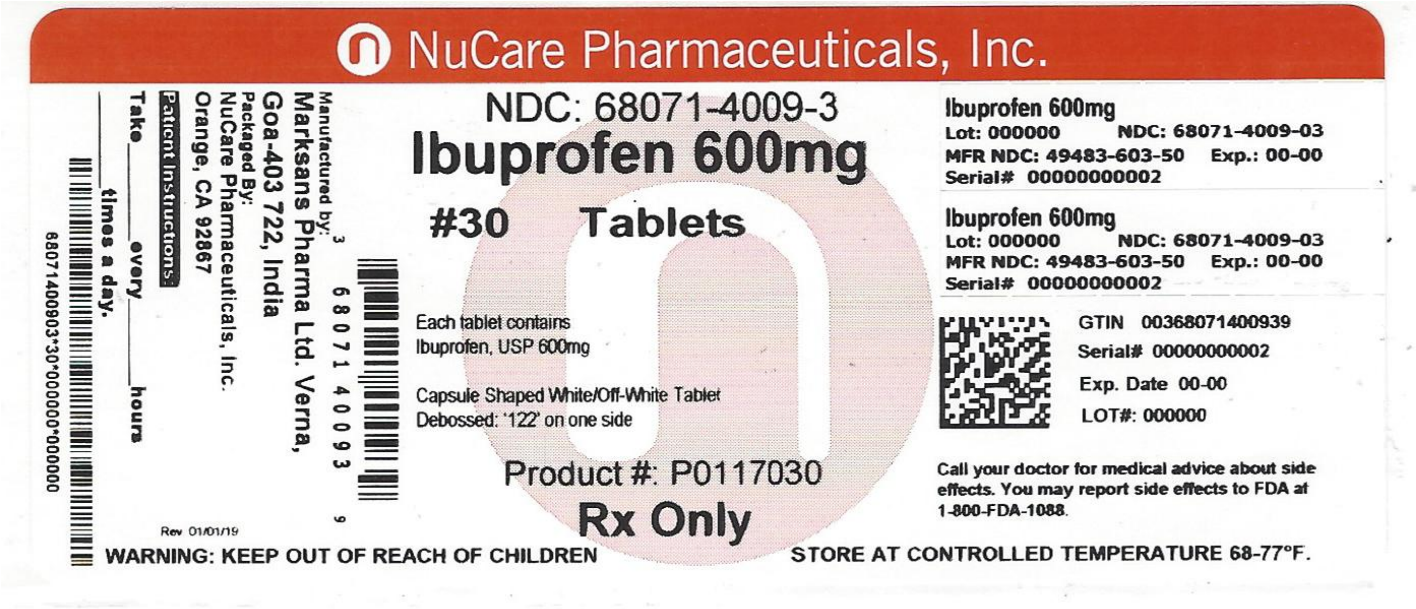
NDC 68071-4009-4 Bottles of 40

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



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)	
CROSCARMELLOSE 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)