

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

HOW SUPPLIED

600mg (white to off white, capsule shaped, biconvex, film coated tablets debossed with '122' on one side and plain on the other side) .

NDC 68071-4243-1 BOTTLES OF 100

600mg Ibuprofen Package Label

NuCare Pharmaceuticals, Inc.

NDC: 68071-4243-1

Ibuprofen 600mg

#100 Tablets

See manufacturer's label for full list of ingredients

Product #: R0284100

Rx Only

Manufactured By: Marksans Pharma Ltd. Verna, Goa-403 722, India

Packaged By: NuCare Pharmaceuticals, Inc. Orange, CA 92867

Take _____ times a day. _____ every _____ hours

Rev 01/01/19

WARNING: KEEP OUT OF REACH OF CHILDREN

STORE AT CONTROLLED TEMPERATURE 68-77°F.

Ibuprofen 600mg
Lot: 000000 NDC: 68071-4243-01
MFR NDC: 49483-603-01 Exp.: 00-00

Ibuprofen 600mg
Lot: 000000 NDC: 68071-4243-01
MFR NDC: 49483-603-01 Exp.: 00-00

GTIN 00368071424317
Serial# 000000000002
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.

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:68071-4243(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:68071-4243-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	01/22/2018	

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	relabel(68071-4243)

Revised: 2/2021

NuCare Pharmaceuticals,Inc.