

IBUPROFEN- ibuprofen tablet, film coated

NuCare Pharmaceuticals, Inc.

HOW SUPPLIED

NDC 68071-3362-0 Bottles of 12

NDC 68071-3362-5 Bottles of 15

NDC 68071-3362-2 Bottles of 20

NDC 68071-3362-1 Bottles of 21

NDC 68071-3362-3 Bottles of 30

NDC 68071-3362-4 Bottles of 40

NDC 68071-3362-6 Bottles of 60

NDC 68071-3362-9 Bottles of 90

800mg Ibuprofen Package Label

**NuCare Pharmaceuticals, Inc.**

NDC: 68071-3362-3
Ibuprofen 800mg
#30 Tablets

Ibuprofen 800mg
Lot: 000000 NDC: 68071-3362-03
MFR NDC: 49483-604-50 Exp.: 00-00
Serial# 00000000002

Ibuprofen 800mg
Lot: 000000 NDC: 68071-3362-03
MFR NDC: 49483-604-50 Exp.: 00-00
Serial# 00000000002

**GTIN 00368071336238**
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
Marksans Pharma Ltd. Verna,
Goa-403 722, India
Packaged By:
NuCare Pharmaceuticals, Inc.
Orange, CA 92667

Take _____ times a day, _____ every _____ hours

68071336203*30*000000*000000

Rev 01/01/19

WARNING: KEEP OUT OF REACH OF CHILDREN

Each tablet contains
Ibuprofen, USP 800mg

Capsule Shaped White/Off-White Tablet
Debossed: '123' on one side

Product #: P0118030
Rx Only

STORE AT CONTROLLED TEMPERATURE 68-77°F.

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type

HUMAN PRESCRIPTION
DRUG

Item Code (Source)

NDC:68071-3362(NDC:49483-
604)

Route of Administration

ORAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

Strength

IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)		IBUPROFEN	800 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	19mm	
Flavor		Imprint Code	123	
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68071-3362-5	15 in 1 BOTTLE; Type 0: Not a Combination Product	07/26/2017	
2	NDC:68071-3362-0	12 in 1 BOTTLE; Type 0: Not a Combination Product	07/26/2017	
3	NDC:68071-3362-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	07/26/2017	
4	NDC:68071-3362-1	21 in 1 BOTTLE; Type 0: Not a Combination Product	07/26/2017	
5	NDC:68071-3362-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	07/26/2017	
6	NDC:68071-3362-4	40 in 1 BOTTLE; Type 0: Not a Combination Product	07/26/2017	
7	NDC:68071-3362-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	07/26/2017	
8	NDC:68071-3362-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	07/26/2017	
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(68071-3362)

Revised: 7/2024

NuCare Pharmaceuticals, Inc.