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

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: 3WQ0SDW1A)	
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.