IBUPROFEN- ibuprofen tablet, film coated
PD-Rx Pharmaceuticals, Inc.

IBUPROFEN 400 MG TABLETS

ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

HOW SUPPLIED

400mg (white to of white, round, biconvex, film coated tablets debossed with '121' on one side and plain on the other side) Bottles of:

NDC 43063-872-06 Bottles of 6

NDC 43063-872-10 Bottles of 10

NDC 43063-872-20 Bottles of 20

NDC 43063-872-30 Bottles of 30

NDC 43063-872-40 Bottles of 40

NDC 43063-872-90 Bottles of 90

NDC 43063-872-82 Bottles of 500

400mg Ibuprofen label

IBUPROFEN

ibuprofen tablet, film coated

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:43063-872(NDC:49483-602)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	400 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	ROUND	**Size**	13mm
Flavor		**Imprint Code**	121
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43063-872-06	6 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/25/2019	
2	NDC:43063-872-10	10 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/29/2019	
3	NDC:43063-872-20	20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/07/2018	
4	NDC:43063-872-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/07/2018	
5	NDC:43063-872-40	40 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/07/2018	
6	NDC:43063-872-90	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/26/2019	
7	NDC:43063-872-82	500 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	03/02/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	12/30/2015	

Labeler - PD-Rx Pharmaceuticals, Inc. (156893695)

Registrant - PD-Rx Pharmaceuticals, Inc. (156893695)

Establishment

Name	Address	ID/FEI	Business Operations
PD-Rx Pharmaceuticals, Inc.		156893695	repack(43063-872)

Revised: 9/2023 PD-Rx Pharmaceuticals, Inc.