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 6, 10, 20, 30, 40, 90 and 500

400mg 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: 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	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: 6/2020 PD-Rx Pharmaceuticals, Inc.