

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

ALCOHOL OR ALCOHOLIC BEVERAGES SHOULD NOT BE CONSUMED WHILE TAKING THIS MEDICATION.

R only

**WARNING: KEEP THIS OUT OF THE REACH OF CHILDREN
DOSAGE and STORAGE: SEE PACKAGE INSERT**

43063-872-90	43063-872-90	43063-872-90
IBUPROFEN	IBUPROFEN	IBUPROFEN
USP	USP	USP
400 MG	400 MG	400 MG
90 TABLETS	90 TABLETS	90 TABLETS
ReOrder # 110794	ReOrder # 110794	ReOrder # 110794
LOT F20A71	LOT F20A71	LOT F20A71
EXP 01/2022	EXP 01/2022	EXP 01/2022

CALL YOUR DOCTOR FOR MEDICAL ADVICE ABOUT SIDE EFFECTS.
YOU MAY REPORT SIDE EFFECTS TO THE FDA AT 1-800-FDA-1088

TAKE ___ TABLET(S) ___ TIMES A DAY WITH FOOD.
TOME ___ TABLETA(S) ___ VECES AL DIA CON COMIDA.

Each TABLET Contains: FILM COATED IBUPROFEN
USP 400 MG



WHITE
121
ROUND

NDC: 43063-872-90



**IBUPROFEN
USP**

400 MG

90 TABLETS



GTIN: 00343063872901
SNO: F20A71000001
EXP: 01/2022
LOT: F20A71

343063872901

4918360320
MANKANS PHARMIA, LTD.
VERINA GDA, 403 722 INDIA

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.