

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

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**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**

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**

<b>43063-872-90</b>	<b>43063-872-90</b>	<b>43063-872-90</b>
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  
ORGANOLEPTIC MARKINGS:  
121  
ROUND

**NDC: 43063-872-90**



**IBUPROFEN  
USP**

**400 MG  
90 TABLETS**



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

4948360260  
MARKSANS PHARMA, LTD.  
VERINA GDA, 403 722 INDIA  
343063872901

## IBUPROFEN

ibuprofen tablet, film coated

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:43063-872(NDC:49483-602)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XY110QM) (IBUPROFEN - UNII:WK2XY110QM)	IBUPROFEN	400 mg

### Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6130)	
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

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	ROUND	<b>Size</b>	13mm
<b>Flavor</b>		<b>Imprint Code</b>	121
<b>Contains</b>			

## 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.