

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

IBUPROFEN 600 MG TABLETS

ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

HOW SUPPLIED

600mg (white to off white, capsule shaped, biconvex, film coated tablets debossed with '122' on one side and plain on the other side) Bottles of 30 NDC 43063-936-30

600 mg label

WARNING KEEP OUT OF
CHILDREN'S REACH

Patient First

DISPENSE IN THIS
TIGHT/LIGHT RESISTANT
CONTAINER

43063-936-30

AFFIX LABEL HERE

DOSAGE AND STORAGE: SEE
PACKAGE INSERT. DISPENSE IN A
TIGHT-LIGHT RESISTANT CONTAINER
AS DEFINED IN THE USP-NF. CALL
YOUR DOCTOR FOR MEDICAL
ADVICE ABOUT SIDE EFFECTS.
YOU MAY REPORT SIDE EFFECTS TO
FDA AT 1-800-FDA-1088

01)10343063936306

(21)Z9Z990002

(17)200831

(10)Z9Z99



IBUPROFEN

USP

600 MG

30 TABLETS

Each Tablet Contains:
IBUPROFEN USP 600 MG

TAKE WITH FOOD



3 43063 93630 6

AFFIX LABEL HERE

MFG BY: Marksans
Pharma Ltd. Plot No. L-82,
L-83, Verna Indl. Estate,
Verna, Goa-403 722, India
603R 1215

Billing Number
49483-0603-50

PACKAGED BY PD-RX
PHARMACEUTICALS, INC
OKLAHOMA CITY, OK
73127

LOT: Z9Z99
EXP:11/2017

RX ONLY

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:43063-936(NDC:49483- 603)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XY110QM) (IBUPROFEN - UNII:WK2XY110QM)	IBUPROFEN	600 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE 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	CAPSULE	Size	18mm
Flavor		Imprint Code	122
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43063-936-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/14/2019	

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(43063-936)

Revised: 9/2023

PD-Rx Pharmaceuticals, Inc.