

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

IBUPROFEN 800 MG TABLETS

ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

800 mg (white to off-white, capsule shaped, biconvex, film-coated□ tablets debossed with ‘123’ on one side and plain on other side)

43063-914-30 Bottles of 30

800 mg label

WARNING KEEP OUT
OF CHILDREN'S
REACH

Patient First

DISPENSE IN THIS
TIGHT/LIGHT RESISTANT
CONTAINER

AFFIX LABEL HERE

DOSAGE AND STORAGE: SEE
PACKAGE OUTSERT. 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

GTIN:00343063914304
SNO: G19E870002
EXP: 201130
LOT: G19E87



43063-914-30

IBUPROFEN

USP

800 MG

30 TABLETS

Each Tablet Contains:
IBUPROFEN USP 800 MG

TAKE WITH FOOD OR MILK.



43063914304

AFFIX LABEL HERE

MFG BY: Marksans
Pharma Ltd. Plot No. L-82,
L-83, Verna Indl. Estate,
Verna, Goa- 403 722, India
Billing Number
49483-604-50
PACKAGED BY PD-RX
PHARMACEUTICALS, INC
OKLAHOMA CITY, OK
73127
LOT: G19E87
EXP:11/2020

RX ONLY

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:43063-914(NDC:49483-604)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XY10QM) (IBUPROFEN - UNII:WK2XY10QM)	IBUPROFEN	800 mg

Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
CELLULOSE, MICROCRYSTALLINE (UNII: OPIR32D61U)	
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	19mm
Flavor		Imprint Code	123
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:43063-914-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/13/2018	

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(43063-914)

Revised: 4/2020

PD-Rx Pharmaceuticals, Inc.