

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) B

NDC 43063-867-10 Bottles of 10

NDC 43063-867-30 Bottles of 30

NDC 43063-867-60 Bottles of 60

NDC 43063-867-01 Bottles of 100

600 mg

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

43063-867-01 IBUPROFEN USP 600 MG 100 TABLETS ReOrder # 110593 LOT F20B02 EXP 07/2021	43063-867-01 IBUPROFEN USP 600 MG 100 TABLETS ReOrder # 110593 LOT F20B02 EXP 07/2021	43063-867-01 IBUPROFEN USP 600 MG 100 TABLETS ReOrder # 110593 LOT F20B02 EXP 07/2021
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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 600 MG



NDC: 43063-867-01



**IBUPROFEN
 USP**

600 MG

100 TABLETS



GTIN: 00343063867013
 SNO: F20B02000002
 EXP: 07/2021
 LOT: F20B02

343063867013
 49483060350
 MARKANS PHARMA, LTD
 VERNAM GOA, 403 722 INDIRA

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:43063-867(NDC:49483-603)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	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:43063-867-10	10 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/05/2018	
2	NDC:43063-867-60	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/05/2018	
3	NDC:43063-867-01	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/11/2018	
4	NDC:43063-867-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/30/2023	

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(43063-867)

Revised: 10/2023

PD-Rx Pharmaceuticals, Inc.