IBUPROFEN- ibuprofen tablet, film coated
REMEDYREPACK INC.

IBUPROFEN 400 MG - 600 MG AND 800 MG TABLETS

ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

Repackaged By / Distributed By: RemedyRepack Inc.

625 Kolter Drive, Indiana, PA 15701

(724) 465-8762

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 100 & 500

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, 50, 100 & 500

600 mg (white to off-white, capsule shaped, biconvex, film-coated tablets debossed with ‘122’ on one side and plain on other side)

NDC: 70518-3313-00

PACKAGING: 30 in 1 BLISTER PACK

Repackaged and Distributed By:

Remedy Repack, Inc.

625 Kolter Dr. Suite #4 Indiana, PA 1-724-465-8762

PRINCIPAL DISPLAY PANEL

DRUG: IBUPROFEN

GENERIC: IBUPROFEN

DOSAGE: TABLET, FILM COATED

ADMINSTRATION: ORAL

NDC: 70518-3313-0

COLOR: white

SHAPE: CAPSULE

SCORE: No score

SIZE: 18 mm

IMPRINT: 122

PACKAGING: 30 in 1 BLISTER PACK

ACTIVE INGREDIENT(S):

- IBUPROFEN 600mg in 1

INACTIVE INGREDIENT(S):

- SILICON DIOXIDE
- CROSCARMELLOSE SODIUM
- MAGNESIUM STEARATE
- CELLULOSE, MICROCRYSTALLINE
- POLYETHYLENE GLYCOL, UNSPECIFIED
- POLYVINYL ALCOHOL
- STARCH, PREGELATINIZED CORN
- TALC
- TITANIUM DIOXIDE

IBUPROFEN

ibuprofen tablet, film coated

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:70518-3313(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)	
CROSCARMELLOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
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			
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:70518-3313-0	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	12/28/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	12/28/2021	

Labeler - REMEDYREPACK INC. (829572556)

Revised: 3/2024 REMEDYREPACK INC.