IBUPROFEN- ibuprofen tablet, film coated
REMEDYREPACK INC.

IBUPROFEN 400 MG - 600 MG AND 800 MG TABLETS

1. ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

Repackaged By / Distributed By: RemedyRepack Inc.

625 Kolter Drive, Indiana, PA 15701

(724) 465-8762

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

NDC: 70518-0005-00

NDC: 70518-0005-01

NDC: 70518-0005-02

NDC: 70518-0005-03

NDC: 70518-0005-04

NDC: 70518-0005-05

NDC: 70518-0005-06

PACKAGING: 21 in 1 BLISTER PACK

PACKAGING: 14 in 1 BLISTER PACK

PACKAGING: 30 in 1 BLISTER PACK

PACKAGING: 10 in 1 BLISTER PACK

PACKAGING: 90 in 1 BOTTLE PLASTIC

PACKAGING: 30 in 1 BOTTLE PLASTIC

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: 70518-0005-0

NDC: 70518-0005-1

NDC: 70518-0005-2

NDC: 70518-0005-3

NDC: 70518-0005-4

NDC: 70518-0005-5

NDC: 70518-0005-6

COLOR: white

SHAPE: CAPSULE

SCORE: No score

SIZE: 19 mm

IMPRINT: 123

PACKAGING: 21 in 1 BLISTER PACK

PACKAGING: 14 in 1 BLISTER PACK

PACKAGING: 30 in 1 BLISTER PACK

PACKAGING: 10 in 1 BLISTER PACK

PACKAGING: 90 in 1 BOTTLE PLASTIC

PACKAGING: 30 in 1 BOTTLE PLASTIC

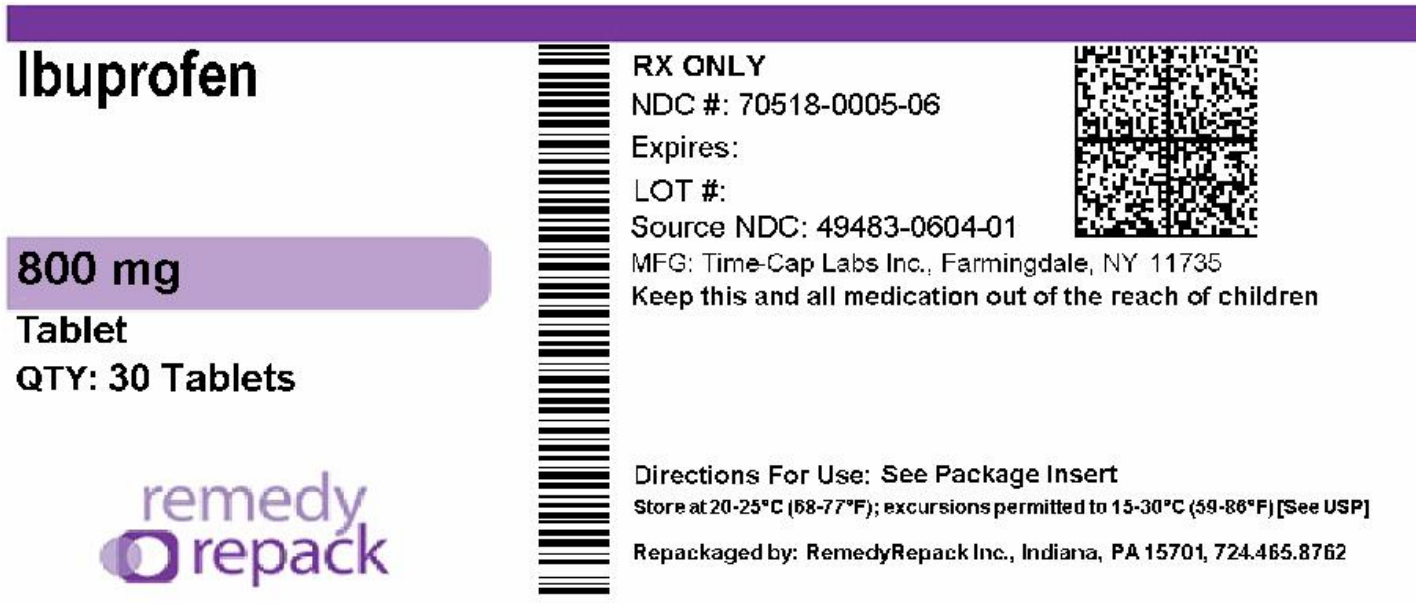
PACKAGING: 30 in 1 BLISTER PACK

ACTIVE INGREDIENT(S):

- IBUPROFEN 800mg 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:70518-0005(NDC:49483-604)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	800 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**	19mm
Flavor		**Imprint Code**	123

Contains

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70518-0005-0	21 in 1 BLISTER PACK; Type 0: Not a Combination Product	11/17/2016	06/04/2018
2	NDC:70518-0005-1	14 in 1 BLISTER PACK; Type 0: Not a Combination Product	12/05/2016	06/01/2018
3	NDC:70518-0005-2	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	12/13/2016	06/11/2018
4	NDC:70518-0005-3	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	01/27/2017	06/11/2018
5	NDC:70518-0005-4	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/06/2017	03/30/2021
6	NDC:70518-0005-5	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/04/2017	06/28/2018
7	NDC:70518-0005-6	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	03/17/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	11/17/2016	

Labeler - REMEDYREPACK INC. (829572556)

Revised: 1/2024 REMEDYREPACK INC.