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
Ibuprofen 400mg, white, round, tablet, imprint: 121
NDC: 70518-2432-00
NDC: 70518-2432-01
PACKAGING: 21 in 1 BLISTER PACK
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: 70518-2432-0

NDC: 70518-2432-1

COLOR: white

SHAPE: ROUND

SCORE: No score

SIZE: 13 mm

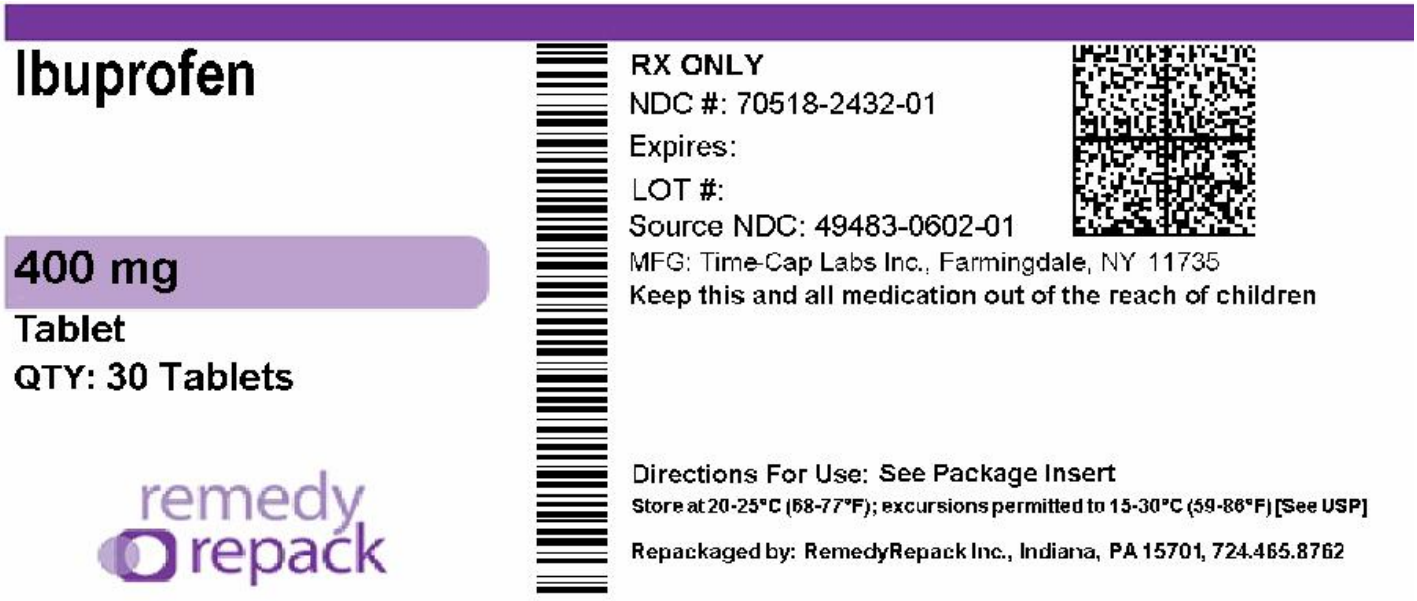
IMPRINT: 121

PACKAGING: 21 in 1 BLISTER PACK

PACKAGING: 30 in 1 BLISTER PACK

ACTIVE INGREDIENT(S):
- IBUPROFEN 400mg 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:70518-2432(NDC:49483-602)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	400 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	ROUND	**Size**	13mm
Flavor		**Imprint Code**	121
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70518-2432-0	21 in 1 BLISTER PACK; Type 0: Not a Combination Product	11/18/2019	11/26/2019
2	NDC:70518-2432-1	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	05/26/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	11/18/2019	

Labeler - REMEDYREPACK INC. (829572556)