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

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

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

PRINCIPAL DISPLAY PANEL

DRUG: IBUPROFEN

GENERIC: IBUPROFEN

DOSAGE: TABLET, FILM COATED

ADMINSTRATION: ORAL

NDC: 70518-2432-0

COLOR: white

SHAPE: ROUND

SCORE: No score

SIZE: 13 mm

IMPRINT: 121

PACKAGING: 21 in 1 BLISTER PACK

ACTIVE INGREDIENT(S):
- IBUPROFEN 400mg in 1

INACTIVE INGREDIENT(S):
- POLYETHYLENE GLYCOLS
- STARCH, PREGELATINIZED CORN
- POLYVINYL ALCOHOL
- TALC
- SILICON DIOXIDE
- CELLULOSE, MICROCRYSTALLINE
- CROSCARMELLOSE SODIUM
- MAGNESIUM STEARATE
- 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: 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	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	

Marketing Information

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

Labeler - REMEDYREPACK INC. (829572556)

Revised: 11/2019 REMEDYREPACK INC.