

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

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

Ibuprofen tablets are available in the following strength:

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

“Preserve in well-closed containers”

Store at 20 ° to 25°C (68°to 77°F) [See USP Controlled Room Temperature].

Rx only

Repackaged and Distributed By:

Remedy Repack, Inc.

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

HOW SUPPLIED

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PRINCIPAL DISPLAY PANEL

DRUG: IBUPROFEN

GENERIC: IBUPROFEN

DOSAGE: TABLET, FILM COATED

ADMINISTRATION: 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 Tablet

MFG NDC: 49483-0604-01

MFG: Time-Cap Labs Inc.,
Farmingdale, NY 11735

800 mg

QTY: 30 Tablets

NDC #: 70518-0005-06

LOT #:

Expires:

Capsule WHITE 123



Repackaged By: RemedyRepack
Inc.,
Indiana, PA 15701, 724.465.8762

RX ONLY

Usual Dosage: See Insert

Keep this and all medication out of
the reach of children

Store at 20-25°C (68-77°F);
excursions permitted to 15-30°C
(59-86°F) [See USP]



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
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: O8232NY35J)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

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	06/04/2019
2	NDC:70518-0005-1	14 in 1 BLISTER PACK; Type 0: Not a Combination Product	12/05/2016	06/01/2019
3	NDC:70518-0005-2	30 in 1 BLISTER PACK; Type 0: Not a Combination Product	12/13/2016	05/11/2019
4	NDC:70518-0005-3	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	01/27/2017	06/11/2019
5	NDC:70518-0005-4	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/06/2017	03/30/2022
6	NDC:70518-0005-5	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	05/04/2017	06/28/2019
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)