

IBUPROFEN- ibuprofen tablet, film coated
medsource pharmaceuticals

IBUPROFEN 800 MG TABLETS

ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

HOW SUPPLIED

400mg (white to off 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)

800 mg 60 count label

IBUPROFEN TABLETS 800MG
 GENERIC FOR MOTRIN

LOT: NDC: 45865-0945-60 EXP: **#60**

UNIT DESCRIPTION: WHITE / OBLONG / 123 / FILM-COATED
 EACH TABLET CONTAINS IBUPROFEN 800MG
 BY: MARKSANS PHARMA LTD. CDA, INDIA

MAY CAUSE DIZZINESS AND OR DROWSINESS. TAKE WITH FOOD TO PREVENT STOMACH UPSET.
 TAKE ___ TABLET(S) ___ TIMES DAILY.

IBUPROFEN TABLETS 800MG
 GENERIC FOR MOTRIN # 60
 LOT: exp: NDC: 45865-0945-60 NFR NDC: 99483-0604-50

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PATIENT LOG CHART CLAIM PEEL HERE

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:45865-945(NDC:49483-604)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XY110QM) (IBUPROFEN - UNII:WK2XY110QM)	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: 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	CAPSULE	Size	19mm
Flavor		Imprint Code	123
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
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1	NDC:45865-945-30	30 in 1 BOTTLE; Type 0: Not a Combination Product	12/12/2018	
2	NDC:45865-945-60	60 in 1 BOTTLE; Type 0: Not a Combination Product	12/12/2018	
Marketing Information				
Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date
ANDA	ANDA090796		12/30/2015	

Labeler - medsource pharmaceuticals (833685915)

Registrant - TIME CAP LABORATORIES, INC (037052099)

Establishment			
Name	Address	ID/FEI	Business Operations
medsource pharmaceuticals		833685915	repack(45865-945)

Revised: 12/2019

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