

IBUPROFEN- ibuprofen tablet, film coated
St. Mary's Medical Park Pharmacy

IBUPROFEN 400 MG - 600 MG AND 800 MG TABLETS

ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

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

Supplied as:

NDC 60760-604-20 BOTTLES OF 20

NDC 60760-604-60 BOTTLES OF 60

NDC 60760-604-90 BOTTLES OF 90

NDC 60760-604-00 BOTTLES OF 100

NDC 60760-604-90

IBUPROFEN
TABLETS, USP
800mg

QTY: 90
LOT# ???????
EXP ??-??
RX# 000000000000
MANUFACTURED BY:
Marksans Pharma Ltd.
Verna, Goa-403 722, India



IBUPROFEN
TABLETS, USP
800mg
QTY: 90
RX# 000000000000
NDC 60760-604-90
LOT# ???????
EXP ??-??

IBUPROFEN
TABLETS, USP
800mg
QTY: 90
RX# 000000000000
NDC 60760-604-90
LOT# ???????
EXP ??-??

IBUPROFEN
TABLETS, USP
800mg
QTY: 90
RX# 000000000000
NDC 60760-604-90
LOT# ???????
EXP ??-??

IBUPROFEN
TABLETS, USP
800mg
QTY: 90
RX# 000000000000
NDC 60760-604-90
LOT# ???????
EXP ??-??

(01) 00360760604901
(21) 000000000000
(17) ???????
(10) ???????



USE AS DIRECTED

PACKAGED BY:
St. Mary's
10860 MAVINEE DR.
ORO VALLEY, AZ 85737
MANAGED PHARMACY PROGRAMS

Rx only STORE AT CONTROLLED ROOM TEMPERATURE 15°- 30° C (59°-86°F)

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	Item Code (Source)	NDC:60760-604(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: 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
1	NDC:60760-604-20	20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/16/2019	
2	NDC:60760-604-60	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/16/2019	
3	NDC:60760-604-90	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/16/2019	
4	NDC:60760-604-00	100 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/14/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	10/16/2019	

Labeler - St. Mary's Medical Park Pharmacy (063050751)**Establishment**

Name	Address	ID/FEI	Business Operations
St. Mary's Medical Park Pharmacy		063050751	relabel(60760-604) , repack(60760-604)

Revised: 4/2020

St. Mary's Medical Park Pharmacy