

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

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**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 60760-604-09 BOTTLES OF 9

NDC 60760-604-20 BOTTLES OF 20

NDC 60760-604-30 BOTTLES OF 30

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**

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:60760-604(NDC:49483-604)
<b>Route of Administration</b>	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**

<b>Color</b>	white	<b>Score</b>	no score
<b>Shape</b>	CAPSULE	<b>Size</b>	19mm
<b>Flavor</b>		<b>Imprint Code</b>	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	
5	NDC:60760-604-09	9 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/10/2020	
6	NDC:60760-604-30	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/08/2020	

**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: 7/2020

St. Mary's Medical Park Pharmacy