

**IBUPROFEN- ibuprofen tablet, film coated**  
**Preferred Pharmaceuticals Inc.**

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**IBUPROFEN 400 MG - 600 MG AND 800 MG TABLETS**

**ibuprofen tablets 400 mg - 600 mg- 800 mg medguide**

Repackaged By: Preferred Pharmaceuticals Inc.

**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 14 NDC 68788-7602-4
- Bottles of 20 NDC 68788-7602-2
- Bottles of 21 NDC 68788-7602-7
- Bottles of 30 NDC 68788-7602-3
- Bottles of 50 NDC 68788-7602-5
- Bottles of 60 NDC 68788-7602-6
- Bottles of 90 NDC 68788-7602-9
- Bottles of 100 NDC 68788-7602-1
- Bottles of 120 NDC 68788-7602-8

Repackaged By: Preferred Pharmaceuticals Inc.

**600mg Label**

# Ibuprofen Tablets 600mg

**PREFERRED**  
Pharmaceuticals, Inc. Anaheim, CA 92807

Generic for: Motrin

Each film-coated tablet contains: Ibuprofen,  
USP 600mg

**Pkg Size:** Exp Date:

Lot#:

Batch#:

Ins:

Mfg: Time-Cap Labs, Inc.; Farmingdale,  
NY

Prod#:

**Warning**

Store at 20°-25°C (68°-77°F). See USP Controlled  
Room Temperature. Rx Only. Keep this and all medication  
out of the reach of children. Tablet is capsule-shaped,  
white, and imprinted with 122.



Directions English

Take \_\_\_ tablet(s)  
every \_\_\_ hours.



Instrucciones Espanol:

Toma \_\_\_ tableta(s)  
cada \_\_\_ horas.

Ibuprofen Tablets 600mg

Qty: Ins:  
Lot#: Bat#:

Prod# (NDC):

Ibuprofen Tablets 600mg

Qty: Ins:  
Lot#: Bat#:

Prod# (NDC):

Ibuprofen Tablets 600mg

Qty: Ins:  
Insurance NDC:  
Lot#: Bat#:

Ibuprofen Tablets 600mg

Qty: Ins:  
Lot#: Bat#:

Prod# (NDC):

Log

Chart

Billing

Patient

## IBUPROFEN

ibuprofen tablet, film coated

### Product Information

<b>Product Type</b>	HUMAN PRESCRIPTION DRUG	<b>Item Code (Source)</b>	NDC:68788-7602(NDC:49483-603)
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XY110QM) (IBUPROFEN - UNII:WK2XY110QM)	IBUPROFEN	600 mg

### Inactive Ingredients

Ingredient Name	Strength
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELOSE SODIUM (UNII: M28OL1HH48)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	
MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WJQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
STARCH, 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>	18mm
<b>Flavor</b>		<b>Imprint Code</b>	122
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-7602-4	14 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2020	
2	NDC:68788-7602-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2020	
3	NDC:68788-7602-7	21 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2020	
4	NDC:68788-7602-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2020	
5	NDC:68788-7602-5	50 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2020	
6	NDC:68788-7602-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2020	
7	NDC:68788-7602-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2020	
8	NDC:68788-7602-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2020	
9	NDC:68788-7602-8	120 in 1 BOTTLE; Type 0: Not a Combination Product	02/14/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	02/14/2020	

**Labeler** - Preferred Pharmaceuticals Inc. (791119022)

**Registrant** - Preferred Pharmaceuticals Inc. (791119022)

## Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals Inc.		791119022	REPACK(68788-7602)

Revised: 2/2020

Preferred Pharmaceuticals Inc.