IBUPROFEN- ibuprofen tablet, film coated
Preferred Pharmaceuticals Inc.

IBUPROFEN 400 MG - 600 MG AND 800 MG TABLETS

ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

Repackaged By: Preferred Pharmaceuticals Inc.

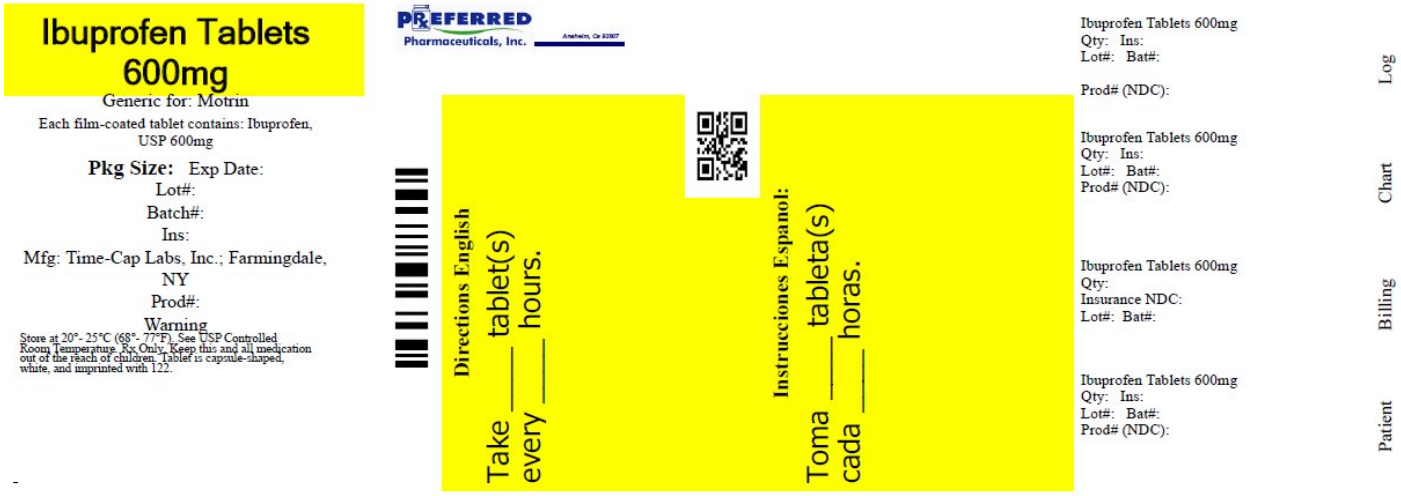
HOW SUPPLIED

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 of100 NDC 68788-7602-1
Bottles of 120 NDC 68788-7602-8

Repackaged By: Preferred Pharmaceuticals Inc.

600mg Label



IBUPROFEN

ibuprofen tablet, film coated

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:68788-7602(NDC:49483-603)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	600 mg

Inactive Ingredients	
Ingredient Name	**Strength**
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
CROSCARMELLOSE 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			
Color	white	**Score**	no score
Shape	CAPSULE	**Size**	18mm
Flavor		**Imprint Code**	122
Contains			

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.