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

HOW SUPPLIED

400mg (white to of white, round, biconvex, film coated tablets debossed with '121' on one side and plain on the other side)

Bottles of 20 NDC 68788-7745-02
Bottles of 28 NDC 68788-7745-08
Bottles of 30 NDC 68788-7745-03
Bottles of 50 NDC 68788-7745-05
Bottles of 60 NDC 68788-7745-06
Bottles of 90 NDC 68788-7745-09
Bottles of 100 NDC 68788-7745-01

Repackaged by Preferred Pharmaceuticals, Inc.

400mg Ibuprofen

IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:68788-7745(NDC:49483-602)
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
IBUPROFEN (UNII: WK2XYI10QM) (IBUPROFEN - UNII:WK2XYI10QM)	IBUPROFEN	400 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: 3WQ0SDW1A)	
POLYVINYL ALCOHOL, UNSPECIFIED (UNII: 532B59J990)	
STARCH, CORN (UNII: O8232NY3SJ)	
TALC (UNII: 7SEV7J4R1U)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	

Product Characteristics

Color	white	**Score**	no score
Shape	ROUND	**Size**	13mm
Flavor		**Imprint Code**	121
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68788-7745-2	20 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
2	NDC:68788-7745-8	28 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
3	NDC:68788-7745-3	30 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
4	NDC:68788-7745-5	50 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
5	NDC:68788-7745-6	60 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
6	NDC:68788-7745-9	90 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	
7	NDC:68788-7745-1	100 in 1 BOTTLE; Type 0: Not a Combination Product	07/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	07/01/2020	

Labeler - Preferred Pharmaceuticals, Inc. (791119022)

Registrant - Preferred Pharmaceuticals, Inc. (791119022)

Establishment

Name	Address	ID/FEI	Business Operations
Preferred Pharmaceuticals, Inc.		791119022	REPACK(68788-7745)

Revised: 5/2024 Preferred Pharmaceuticals, Inc.