IBUPROFEN- ibuprofen tablet, film coated
Northwind Pharmaceuticals

IBUPROFEN 600 MG TABLETS

ibuprofen tablets 400 mg - 600 mg- 800 mg medguide

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 20 NDC: 51655-474-20, 30 NDC: 51655-474-52, 40 NDC: 51655-474-51, 60 NDC: 51655-474-25, and 90 NDC: 51655-474-90.

600 mg 30 count label

NDC: 51655-474-52

IBUPROFEN

ibuprofen tablet, film coated

Product Information			
Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:51655-474(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)	
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
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**	18mm
Flavor		**Imprint Code**	122

Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51655-474-20	20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/19/2020	
2	NDC:51655-474-25	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/13/2020	
3	NDC:51655-474-51	40 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/08/2021	
4	NDC:51655-474-52	30 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/14/2021	
5	NDC:51655-474-26	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/21/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	11/13/2020	

Labeler - Northwind Pharmaceuticals (036986393)

Registrant - Northwind Pharmaceuticals (036986393)

Establishment

Name	Address	ID/FEI	Business Operations
Northwind Pharmaceuticals		036986393	repack(51655-474)

Revised: 1/2023 Northwind Pharmaceuticals