IBUPROFEN- ibuprofen tablet, film coated
Northwind Pharmaceuticals

IBUPROFEN 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)

NDC: 51655-275-54 15 Bottles

NDC: 51655-275-20 20 Bottles

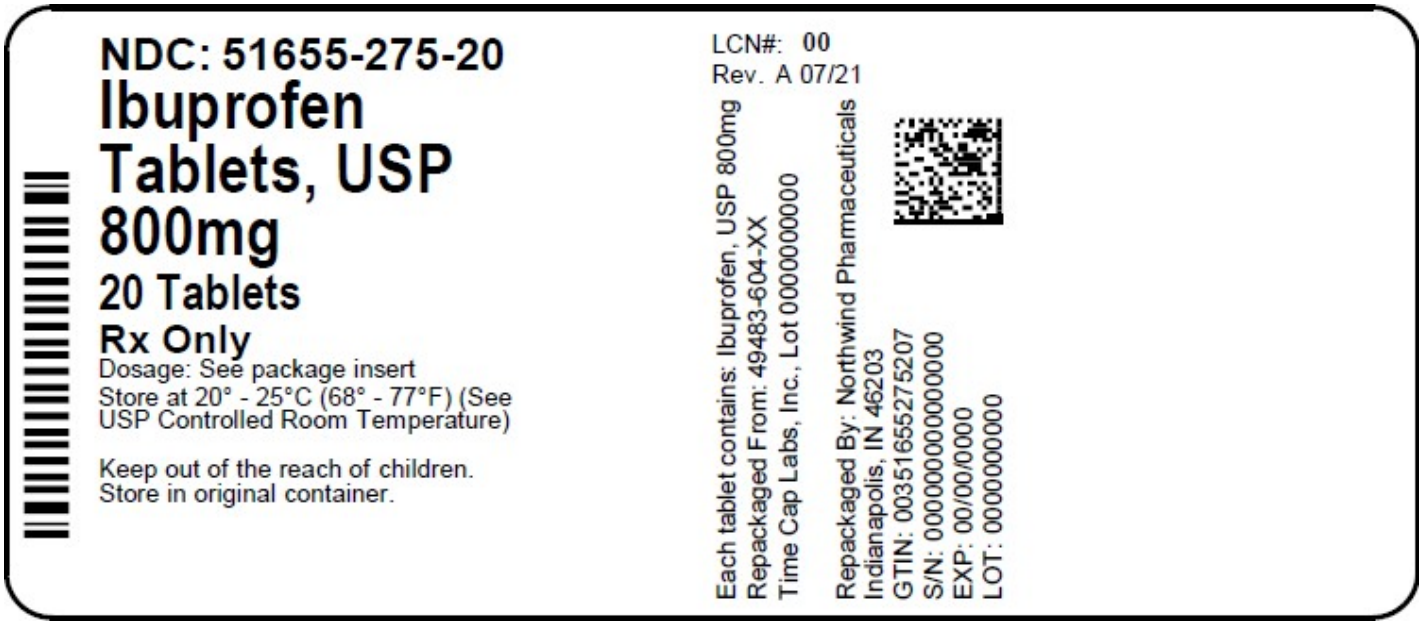
NDC: 51655-275-21 21 Bottles

NDC: 51655-275-25 60 Bottles

NDC: 51655-275-26 90 Botlles

800 mg 20 count label

NDC: 51655-275-20



IBUPROFEN

ibuprofen tablet, film coated

Product Information

Product Type	HUMAN PRESCRIPTION DRUG	**Item Code (Source)**	NDC:51655-275(NDC:49483-604)
Route of Administration	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: 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**	19mm
Flavor		**Imprint Code**	123

Contains	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51655-275-20	20 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/09/2021	
2	NDC:51655-275-21	21 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/29/2020	
3	NDC:51655-275-25	60 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/17/2021	
4	NDC:51655-275-26	90 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/16/2020	
5	NDC:51655-275-54	15 in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/21/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA090796	06/16/2020	

Labeler - Northwind Pharmaceuticals (036986393)

Registrant - Northwind Pharmaceuticals (036986393)

Establishment

Name	Address	ID/FEI	Business Operations
Northwind Pharmaceuticals		036986393	repack(51655-275)

Revised: 1/2023 Northwind Pharmaceuticals