Drug Details

Drug Name(s)	**IBUPROFEN**
FDA Application No.	**(ANDA) 203599**
Active Ingredient(s)	**IBUPROFEN**
Company	**SOFGEN PHARMS**
Original Approval or Tentative Approval Date	**September 7, 2016**

- **Other OTC Drugs with the same Active Ingredient, Strength and Dosage Form/Route**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #203599
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
IBUPROFEN	IBUPROFEN	EQ 200MG FREE ACID AND POTASSIUM SALT	CAPSULE;ORAL	Over-the-counter	No	None

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