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## Drug Details

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

- [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:**

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

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