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Drug Details

Drug Name(s)	NAPROXEN SODIUM
FDA Application No.	(ANDA) 205497
Active Ingredient(s)	NAPROXEN SODIUM
Company	AUROBINDO PHARMA LTD
Original Approval or Tentative Approval Date	March 18, 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) #205497

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
NAPROXEN SODIUM	NAPROXEN SODIUM	EQ 200MG BASE	TABLET;ORAL	Over-the-counter	No None

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