Drugs@FDA
FDA Approved Drug Products

Drug Details

Drug Name(s)	**BUPIVACAINE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 207183**
Active Ingredient(s)	**BUPIVACAINE HYDROCHLORIDE**
Company	**AUROBINDO PHARMA LTD**
Original Approval or Tentative Approval Date	**May 13, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #207183
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
BUPIVACAINE HYDROCHLORIDE	BUPIVACAINE HYDROCHLORIDE	0.25%	INJECTABLE;INJECTION	Prescription	No	AP
BUPIVACAINE HYDROCHLORIDE	BUPIVACAINE HYDROCHLORIDE	0.5%	INJECTABLE;INJECTION	Prescription	No	AP

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