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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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