Drug Details

Drug Name(s)	**ROPIVACAINE HYDROCHLORIDE**
FDA Application No.	**(ANDA) 205612**
Active Ingredient(s)	**ROPIVACAINE HYDROCHLORIDE**
Company	**AUROBINDO PHARMA LTD**
Original Approval or Tentative Approval Date	**July 13, 2016**

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

Products on Application (ANDA) #205612
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
ROPIVACAINE HYDROCHLORIDE	ROPIVACAINE HYDROCHLORIDE	40MG/20ML (2MG/ML)	SOLUTION;INJECTION	Prescription	No	AP
ROPIVACAINE HYDROCHLORIDE	ROPIVACAINE HYDROCHLORIDE	200MG/100ML (2MG/ML)	SOLUTION;INJECTION	Prescription	No	AP
ROPIVACAINE HYDROCHLORIDE	ROPIVACAINE HYDROCHLORIDE	100MG/20ML (5MG/ML)	SOLUTION;INJECTION	Prescription	No	AP
ROPIVACAINE HYDROCHLORIDE	ROPIVACAINE HYDROCHLORIDE	150MG/30ML (5MG/ML)	SOLUTION;INJECTION	Prescription	No	AP
ROPIVACAINE HYDROCHLORIDE	ROPIVACAINE HYDROCHLORIDE	150MG/20ML (7.5MG/ML)	SOLUTION;INJECTION	Prescription	No	AP
ROPIVACAINE HYDROCHLORIDE	ROPIVACAINE HYDROCHLORIDE	100MG/10ML (10MG/ML)	SOLUTION;INJECTION	Prescription	No	AP
ROPIVACAINE HYDROCHLORIDE	ROPIVACAINE HYDROCHLORIDE	200MG/20ML (10MG/ML)	SOLUTION;INJECTION	Prescription	No	AP

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