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Drug Details

Drug Name(s)	ROIIVACAINE HYDROCHLORIDE
FDA Application No.	(ANDA) 205612
Active Ingredient(s)	ROIIVACAINE 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	RLDTE Code
ROIIVACAINE HYDROCHLORIDE	ROIIVACAINE HYDROCHLORIDE	40MG/20ML (2MG/ML)	SOLUTION;INJECTION	Prescription No	AP
ROIIVACAINE HYDROCHLORIDE	ROIIVACAINE HYDROCHLORIDE	200MG/100ML (2MG/ML)	SOLUTION;INJECTION	Prescription No	AP
ROIIVACAINE HYDROCHLORIDE	ROIIVACAINE HYDROCHLORIDE	100MG/20ML (5MG/ML)	SOLUTION;INJECTION	Prescription No	AP
ROIIVACAINE HYDROCHLORIDE	ROIIVACAINE HYDROCHLORIDE	150MG/30ML (5MG/ML)	SOLUTION;INJECTION	Prescription No	AP
ROIIVACAINE HYDROCHLORIDE	ROIIVACAINE HYDROCHLORIDE	150MG/20ML (7.5MG/ML)	SOLUTION;INJECTION	Prescription No	AP
ROIIVACAINE HYDROCHLORIDE	ROIIVACAINE HYDROCHLORIDE	100MG/10ML (10MG/ML)	SOLUTION;INJECTION	Prescription No	AP
ROIIVACAINE HYDROCHLORIDE	ROIIVACAINE HYDROCHLORIDE	200MG/20ML (10MG/ML)	SOLUTION;INJECTION	Prescription No	AP

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