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Drug Details

Drug Name(s)	AMIODARONE HYDROCHLORIDE
FDA Application No.	(ANDA) 204742
Active Ingredient(s)	AMIODARONE HYDROCHLORIDE
Company	AUROBINDO PHARMA LTD
Original Approval or Tentative Approval Date	June 3, 2016

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- [Approval History, Letters, Reviews, and Related Documents](#)
- **Labels are not available**

Products on Application (ANDA) #204742

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
AMIODARONE HYDROCHLORIDE	AMIODARONE HYDROCHLORIDE	200MG	TABLET;ORAL	Prescription	No	AB

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