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Drug Details

Drug Name(s)	DEXAMETHASONE SODIUM PHOSPHATE
FDA Application No.	(ANDA) 206781
Active Ingredient(s)	DEXAMETHASONE SODIUM PHOSPHATE
Company	AUROBINDO PHARMA LTD
Original Approval or Tentative Approval Date	December 1, 2015

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

Products on Application (ANDA) #206781

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
DEXAMETHASONE SODIUM PHOSPHATE	DEXAMETHASONE SODIUM PHOSPHATE	EQ 4MG PHOSPHATE/ML	INJECTABLE;INJECTION	Prescription No	AP

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