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Drug Details

Drug Name(s)	CAFFEINE CITRATE
FDA Application No.	(ANDA) 205013
Active Ingredient(s)	CAFFEINE CITRATE
Company	AUROBINDO PHARMA LTD
Original Approval or Tentative Approval Date	September 22, 2015

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

Products on Application (ANDA) #205013

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
CAFFEINE CITRATE	CAFFEINE CITRATE	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	SOLUTION;INTRAVENOUS	Prescription No	AP

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