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Drug Details

Drug Name(s)	AMPICILLIN SODIUM
FDA Application No.	(ANDA) 202864
Active Ingredient(s)	AMPICILLIN SODIUM
Company	HOSPIRA INC
Original Approval or Tentative Approval Date	September 4, 2015

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

Products on Application (ANDA) #202864

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
AMPICILLIN SODIUM	AMPICILLIN SODIUM	EQ 250MG BASE/VIAL	INJECTABLE;INJECTION	Prescription	No	AP
AMPICILLIN SODIUM	AMPICILLIN SODIUM	EQ 500MG BASE/VIAL	INJECTABLE;INJECTION	Prescription	No	AP
AMPICILLIN SODIUM	AMPICILLIN SODIUM	EQ 1GM BASE/VIAL	INJECTABLE;INJECTION	Prescription	No	AP
AMPICILLIN SODIUM	AMPICILLIN SODIUM	EQ 2GM BASE/VIAL	INJECTABLE;INJECTION	Prescription	No	AP

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