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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