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Drug Details

Drug Name(s)	NEVIRAPINE
FDA Application No.	(ANDA) 204621
Active Ingredient(s)	NEVIRAPINE
Company	ALVOGEN PINE BROOK
Original Approval or Tentative Approval Date	November 9, 2015

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

Products on Application (ANDA) #204621

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD TE Code
NEVIRAPINE	NEVIRAPINE	400MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No AB
NEVIRAPINE	NEVIRAPINE	100MG	TABLET, EXTENDED RELEASE;ORAL	None (Tentative Approval)	No None

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