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Drug Details

Drug Name(s)	NEVIRAPINE
FDA Application No.	(ANDA) 206271
Active Ingredient(s)	NEVIRAPINE
Company	MYLAN PHARMS INC
Original Approval or Tentative Approval Date	November 9, 2015

- [Therapeutic Equivalents](#)
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- **Labels are not available**

Products on Application (ANDA) #206271

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	100MG	TABLET, EXTENDED RELEASE;ORAL	Prescription	No	AB

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