Drug Details

Drug Name(s)	**PROBUPHINE**
FDA Application No.	**(NDA) 204442**
Active Ingredient(s)	**BUPRENORPHINE HYDROCHLORIDE**
Company	**TITAN PHARMS**
Original Approval or Tentative Approval Date	**May 26, 2016**

- **There are no Therapeutic Equivalents**
- **Label Information**
- **Approval History, Letters, Reviews, and Related Documents**

Products on Application (NDA) #204442
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
PROBUPHINE	BUPRENORPHINE HYDROCHLORIDE	80MG	IMPLANT;IMPLANTATION	Prescription	TBD[11]	TBD[12]

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