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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](#)
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- [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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